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**A global synthesis of the effects of diversified farming systems on arthropod diversity within fields and across agricultural landscapes**

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**A global synthesis of the effects of diversified farming systems on arthropod diversity within fields and across agricultural landscapes**

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Complete List of Authors:	<p>Lichtenberg, Elinor; University of Arizona, Department of Ecology and Evolutionary Biology; Washington State University, Department of Entomology</p> <p>Kennedy, Christina; The Nature Conservancy, Global Conservation Lands Program</p> <p>Kremen, Claire; University of California Berkeley, Department of Environmental Sciences, Policy and Management</p> <p>Batary, Peter; Georg-August University, Agroecology</p> <p>Berendse, Frank; Wageningen Universiteit, Nature Conservation and Plant Ecology Group</p> <p>Bommarco, Riccardo; Swedish University of Agricultural Sciences, Department of Ecology</p> <p>Bosque-Pérez, Nilsa; University of Idaho, Department of Plant, Soil and Entomological Sciences</p> <p>Carvalho, Luísa; Universidade de Brasília, Campus Universitário Darcy, Departamento de Ecologia; Faculdade de Ciências da Universidade de Lisboa, Center for Ecology, Evolution and Environmental Changes (CE3C)</p> <p>Snyder, William; Washington State University, Department of Entomology</p> <p>Williams, Neal; UC Davis, Department of Entomology</p> <p>Winfrey, Rachael; Rutgers University, Department of Ecology, Evolution and Natural Resources</p> <p>Åström, Sandra; Norwegian Institute for Nature Research - NINA,</p> <p>Benjamin, Faye; Rutgers University, Department of Ecology, Evolution and Natural Resources</p> <p>Brittain, Claire; UC Davis, Department of Entomology</p> <p>Chaplin-Kramer, Rebecca; Stanford University,</p> <p>Clough, Yann; Lund University, Centre for Environmental and Climate Research</p> <p>Connelly, Heather; Cornell University, Department of Entomology</p> <p>Danforth, Bryan; Cornell University, Department of Entomology</p> <p>Diekötter, Tim; Kiel University, Department of Landscape Ecology</p> <p>Eigenbrode, Sanford; University of Idaho, Department of Plant, Soil and Entomological Sciences</p> <p>Ekroos, Johan; Lund University, Centre for Environmental and Climate Research</p> <p>Elle, Elizabeth; Simon Fraser University, Department of Biological Sciences</p>

	<p>Freitas, Breno; Universidade Federal do Ceará, Departamento de Zootecnia</p> <p>Fukuda, Yuki; University of Otago, Centres for the Study of Agriculture Food and Environment</p> <p>Gaines-Day, Hannah; University of Wisconsin Madison, Department of Entomology</p> <p>Gratton, Claudio; University of Wisconsin Madison, Department of Entomology</p> <p>Holzschuh, Andrea; University of Würzburg, Department of Animal Ecology and Tropical Biology</p> <p>Isaacs, Rufus; Michigan State University, Department of Entomology</p> <p>Isaia, Marco; Università degli Studi di Torino Scuola di Scienze della Natura, Life Sciences and Systems Biology</p> <p>Jha, Shalene; University of Texas at Austin, Department of Integrative Biology</p> <p>Jonason, Dennis; Stockholm University, Department of Physical Geography</p> <p>Jones, Vincent; Tree Fruit Research and Extension Center, Washington State University, Department of Entomology</p> <p>Klatt, Björn; Georg-August University, Agroecology; Lund University, Centre for Environmental and Climate Research; Lund University, Department of Biology</p> <p>Klein, Alexandra-Maria; University of Freiburg, Nature Conservation and Landscape Ecology</p> <p>Krauss, Jochen; Julius-Maximilians-Universität Würzburg Institut für deutsche Philologie, Department of Animal Ecology and Tropical Biology</p> <p>Letourneau, Deborah; University of California, Santa Cruz, Department of Environmental Studies</p> <p>MacFadyen, Sarina; CSIRO</p> <p>Mallinger, Rachel; University of Wisconsin, Department of Entomology</p> <p>Martinez, Eliana; CORPOICA, Centro de Investigación Obonuco</p> <p>Memmott, Jane; University of Bristol, School of Biological Sciences</p> <p>Morandin, Lora; Pollinator Partnership Canada</p> <p>Neame, Lisa; Alberta Environment and Parks, Regional Planning Branch</p> <p>Otieno, Mark; Embu University College, Department of Agricultural Resource Management</p> <p>Park, Mia; Cornell University, Department of Entomology; University of North Dakota, Department of Humanities &amp; Integrated Studies</p> <p>Pfiffner, Lukas; Forschungsinstitut für biologischen Landbau, Department of Crop Science</p> <p>Pocock, Michael; Centre for Ecology &amp; Hydrology</p> <p>Ponce, Carlos; Museo Nacional de Ciencias Naturales, Department of Evolutionary Ecology</p> <p>Potts, Simon; University of Reading, Centre for Agri-Environmental Research, School of Agriculture, Policy and Development</p> <p>Poveda, Katja; Cornell University, Department of Entomology</p> <p>Ramos, Mariangie; University of Puerto Rico at Utuado, Department of Agricultural Technology</p> <p>Rosenheim, Jay; UC Davis, Department of Entomology</p> <p>Rundlöf, Maj; Lunds Universitet, Department of Biology</p> <p>Sardinas, Hilary; University of California Berkeley, Department of Environmental Sciences, Policy and Management</p> <p>Saunders, Manu; Charles Sturt University, Institute for Land Water and Society</p> <p>Schon, Nicole; AgResearch Ltd Lincoln Research Centre</p> <p>Sciligo, Amber; University of California Berkeley, Department of Environmental Sciences, Policy and Management</p> <p>Sidhu, C.; University of California Cooperative Extension, San Mateo &amp; San Francisco Counties</p> <p>Steffan-Dewenter, Ingolf; University of Würzburg, Department of Animal Ecology and Tropical Biology</p> <p>Tscharntke, Teja; Georg-August University, Agroecology</p> <p>Vesely, Milan; Univerzita Palackého v Olomouci, Department of Zoology,</p>
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	<p>Faculty of Science  Weisser, Wolfgang; Technical University of Munich, Terrestrial Ecology Research Group, Department for Ecology and Ecosystem Management  Wilson, Julianna; Michigan State University, Department of Entomology  Crowder, David; Washington State University, Department of Entomology</p>
Keywords:	<p>agricultural management schemes, arthropod diversity, functional groups, landscape complexity, meta-analysis, evenness, biodiversity, organic farming</p>
Abstract:	<p>Agricultural intensification is a leading cause of global biodiversity loss, which can reduce the provisioning of ecosystem services in managed ecosystems. Organic farming and plant diversification are farm management schemes that may mitigate potential ecological harm by increasing species richness and boosting related ecosystem services to agroecosystems. What remains unclear is the extent to which farm management schemes affect biodiversity components other than species richness, and whether impacts differ across spatial scales and landscape contexts. Using a global meta-dataset, we quantified the effects of organic farming and plant diversification on abundance, local diversity (communities within fields), and regional diversity (communities across fields) of arthropod pollinators, predators, herbivores, and detritivores. Both organic farming and higher in-field plant diversity enhanced arthropod abundance, particularly for rare taxa. This resulted in increased richness but decreased evenness. While these responses were stronger at local relative to regional scales, richness and abundance increased at both scales, and richness on farms embedded in complex relative to simple landscapes. Overall, both organic farming and in-field plant diversification exerted the strongest effects on pollinators and predators, suggesting these management schemes can facilitate ecosystem service providers without augmenting herbivore (pest) populations. Our results suggest that organic farming and plant diversification promote diverse arthropod meta-communities that may provide temporal and spatial stability of ecosystem service provisioning. Conserving diverse plant and arthropod communities in farming systems therefore requires sustainable practices that operate both within fields and across landscapes.</p>

A global synthesis of the effects of diversified farming systems on arthropod diversity within fields and across agricultural landscapes

**Running head:** Effects of diversified farming on arthropods

Elinor M. Lichtenberg<sup>1,2</sup>, Christina M. Kennedy<sup>3</sup>, Claire Kremen<sup>4</sup>, Péter Batáry<sup>5</sup>, Frank Berendse<sup>6</sup>, Riccardo Bommarco<sup>7</sup>, Nilsa A. Bosque-Pérez<sup>8</sup>, Luísa G. Carvalheiro<sup>9,10</sup>, William E. Snyder<sup>1</sup>, Neal M. Williams<sup>11</sup>, Rachael Winfree<sup>12</sup>, Sandra Åström<sup>13</sup>, Faye Benjamin<sup>12</sup>, Claire Brittain<sup>11</sup>, Rebecca Chaplin-Kramer<sup>14</sup>, Yann Clough<sup>15</sup>, Heather Connelly<sup>16</sup>, Bryan Danforth<sup>16</sup>, Tim Diekötter<sup>17</sup>, Sanford D. Eigenbrode<sup>8</sup>, Johan Ekroos<sup>15</sup>, Elizabeth Elle<sup>18</sup>, Breno M. Freitas<sup>19</sup>, Yuki Fukuda<sup>20</sup>, Hannah R. Gaines-Day<sup>21</sup>, Claudio Gratton<sup>21</sup>, Andrea Holzschuh<sup>22</sup>, Rufus Isaacs<sup>23</sup>, Marco Isaia<sup>24</sup>, Shalene Jha<sup>25</sup>, Dennis Jonason<sup>26</sup>, Vincent P. Jones<sup>27</sup>, Björn Klatt<sup>5,15,28</sup>, Alexandra-Maria Klein<sup>29</sup>, Jochen Krauss<sup>22</sup>, Deborah K. Letourneau<sup>30</sup>, Sarina Macfadyen<sup>31</sup>, Rachel E. Mallinger<sup>21</sup>, Emily A. Martin<sup>22</sup>, Eliana Martinez<sup>32</sup>, Jane Memmott<sup>33</sup>, Lora Morandin<sup>34</sup>, Lisa Neame<sup>35</sup>, Mark Otieno<sup>36</sup>, Mia G. Park<sup>16,37</sup>, Lukas Pfiffner<sup>38</sup>, Michael Pocock<sup>39</sup>, Carlos Ponce<sup>40</sup>, Simon G. Potts<sup>41</sup>, Katja Poveda<sup>16</sup>, Mariangie Ramos<sup>42</sup>, Jay A. Rosenheim<sup>11</sup>, Maj Rundlöf<sup>28</sup>, Hilary Sardiñas<sup>4</sup>, Manu E. Saunders<sup>43</sup>, Nicole L. Schon<sup>44</sup>, Amber R. Sciligo<sup>4</sup>, C. Sheena Sidhu<sup>45</sup>, Ingolf Steffan-Dewenter<sup>22</sup>, Teja Tscharntke<sup>5</sup>, Milan Veselý<sup>46</sup>, Wolfgang W. Weisser<sup>47</sup>, Julianna K. Wilson<sup>23</sup>, David W. Crowder<sup>1</sup>

(1) Department of Entomology, Washington State University, Pullman, WA, USA (Elinor M. Lichtenberg, elinor@email.arizona.edu; William E. Snyder, wesnyder@wsu.edu; David W. Crowder, dcrowder@wsu.edu)

(2) Department of Ecology & Evolutionary Biology, The University of Arizona, Tucson, AZ, USA

(3) Global Conservation Lands Program, The Nature Conservancy, Fort Collins, CO, USA (ckennedy@tnc.org)

(4) Department of Environmental Sciences, Policy and Management, University of California, Berkeley, CA, USA (Claire Kremen, ckremen@berkeley.edu; Hilary Sardiñas, hsardinas@berkeley.edu; Amber Sciligo, amber.sciligo@berkeley.edu)

(5) Agroecology, University of Göttingen, Grisebachstrasse 6, 37077 Göttingen, Germany (Péter Batáry, pbatary@gmail.com; Bjorn Klatt, klattbk@googlemail.com; Teja Tscharntke, ttschar@gwdg.de)

(6) Nature Conservation and Plant Ecology Group, Wageningen University, PO Box 47, 6700 AA Wageningen, the Netherlands, (frank.berendse@wur.nl)

(7) Swedish University of Agricultural Sciences, Department of Ecology, 75007 Uppsala, Sweden (Riccardo.Bommarco@slu.se)

(8) Department of Plant, Soil and Entomological Sciences, University of Idaho, Moscow, ID, USA (Nilsa Bosque-Pérez, nbosque@uidaho.edu; Sanford D. Eigenbrode, sanforde@uidaho.edu)

(9) Departamento de Ecologia, Universidade de Brasília, Campus Universitário Darcy

- 43 Ribeiro, Brasília - DF, 70910-900, Brazil (lgcarvalho@gmail.com)
- 44 (10) Center for Ecology, Evolution and Environmental Changes (CE3C), Faculdade de  
45 Ciencias da Universidade de Lisboa, 1749-016 Lisboa, Portugal
- 46 (11) Department of Entomology and Nematology, University of California, Davis, Davis,  
47 CA, USA (Neal M. Williams, nmwilliams@ucdavis.edu; Claire Brittain,  
48 claireabrittain@gmail.com; Jay A. Rosenheim, jarozenheim@ucdavis.edu)
- 49 (12) Department of Ecology, Evolution and Natural Resources, Rutgers University, New  
50 Brunswick, NJ, USA (Rachael Winfree, rwinfree@rutgers.edu; Faye Benjamin,  
51 faye.benjamin@gmail.com)
- 52 (13) Norwegian Institute for Nature Research (NINA), PO box 5685 Sluppen, NO-7485  
53 Trondheim, Norway (sandra.astrom@nina.no)
- 54 (14) Natural Capital Project, Stanford University, Stanford, CA, USA  
55 (bchaplin@stanford.edu)
- 56 (15) Centre for Environmental and Climate Research, Lund University, Lund, Sweden  
57 (Yann Clough, yann.clough@cec.lu.se; Johan Ekroos, johan.ekroos@cec.lu.se)
- 58 (16) Department of Entomology, Cornell University, Ithaca, NY, USA (Heather Connelly,  
59 hlc66@cornell.edu; Bryan Danforth, bnd1@cornell.edu; Mia G. Park, mia.park@und.edu;  
60 Katja Poveda, kap235@cornell.edu)
- 61 (17) Department of Landscape Ecology, Kiel University, Kiel, Germany  
62 (tdiekoetter@ecology.uni-kiel.de)
- 63 (18) Department of Biological Sciences, Simon Fraser University, Burnaby, BC, Canada,  
64 Canada (eelle@sfu.ca)
- 65 (19) Departamento de Zootecnia, Universidade Federal do Ceará, 60.356-000 Fortaleza,  
66 CE, Brazil (freitas@ufc.br).
- 67 (20) Centres for the Study of Agriculture Food and Environment, University of Otago,  
68 Dunedin 9054, New Zealand (yuki\_fukuda@hotmail.com)
- 69 (21) Department of Entomology, University of Wisconsin-Madison, Madison, WI USA  
70 (Hanna R. Gaines-Day, hgaines@wisc.edu; Claudio Gratton, cgratton@wisc.edu; Rachel E.  
71 Mallinger, rachel.mallinger@gmail.com)
- 72 (22) Department of Animal Ecology and Tropical Biology, Biocenter, University of  
73 Würzburg, Am Hubland, 97074 Würzburg, Germany (Andrea Holzschuh,  
74 andrea.holzschuh@uni-wuerzburg.de; Jochen Krauss, j.krauss@uni-wuerzburg.de; Emily A.  
75 Martin, emily.martin@uni-wuerzburg.de; Ingolf Steffan-Dewenter, ingolf.steffan@uni-  
76 wuerzburg.de)
- 77 (23) Department of Entomology, Michigan State University, East Lansing, MI, USA  
78 (isaacsr@msu.edu, jkwilson@msu.edu)
- 79 (24) Department of Life Sciences and Systems Biology, University of Torino, Italy  
80 (marco.isaia@unito.it)
- 81 (25) Department of Integrative Biology, University of Texas at Austin, Austin, TX, USA

- 82 (sjha@austin.utexas.edu)
- 83 (26) Department of Physical Geography, Stockholm University, Stockholm, Sweden  
84 (dennis.jonason@natgeo.su.se)
- 85 (27) Department of Entomology, Tree Fruit Research and Extension Center, Washington  
86 State University, Wenatchee, WA, USA (vpjones@wsu.edu)
- 87 (28) Department of Biology, Lund University, Lund, Sweden (maj.rundlof@biol.lu.se)
- 88 (29) Nature Conservation and Landscape Ecology, Faculty of Environment and Natural  
89 Resources, University of Freiburg, Freiburg, Germany (alexandra.klein@nature.uni-  
90 freiburg.de)
- 91 (30) Department of Environmental Studies, University of California, Santa Cruz, CA,  
92 USA (dletour@ucsc.edu)
- 93 (31) CSIRO, Clunies Ross St, Acton ACT 2601, Australia, (sarina.macfadyen@csiro.au)
- 94 (32) CORPOICA, Centro de Investigación Obonuco. Km 5 vía Obonuco - Pasto, Nariño,  
95 Colombia (Emartinezp@corpoica.org.co)
- 96 (33) School of Biological Sciences, University of Bristol, Bristol, BS8 1TQ, UK  
97 (Jane.Memmott@bristol.ac.uk)
- 98 (34) Pollinator Partnership Canada, Victoria, BC, Canada (lm@pollinator.org)
- 99 (35) Alberta Environment and Parks, Regional Planning Branch, Edmonton, Alberta,  
100 Canada (lisa.neame@gov.ab.ca)
- 101 (36) Department of Agricultural Resource Management, Embu University College, Embu,  
102 Kenya (mmarkotieno@gmail.com)
- 103 (37) Department of Humanities & Integrated Studies, University of North Dakota, Grand  
104 Forks, ND, USA
- 105 (38) Department of Crop Science, Research Institute of Organic Agriculture, Frick,  
106 Switzerland (lukas.pfiffner@fibl.org)
- 107 (39) NERC Centre for Ecology & Hydrology, Crowmarsh Gifford, Wallingford,  
108 Oxfordshire, United Kingdom (michael.pocock@ceh.ac.uk)
- 109 (40) Department of Evolutionary Ecology, Museo Nacional de Ciencias Naturales, CSIC,  
110 Madrid, Spain (carlosp@mncn.csic.es)
- 111 (41) Centre for Agri-Environmental Research, School of Agriculture, Policy and  
112 Development, University of Reading, Reading, RG66AR, UK (s.g.potts@reading.ac.uk)
- 113 (42) Department of Agricultural Technology, University of Puerto Rico at Utuado,  
114 Utuado, PR, USA (mariangie.ramos@upr.edu)
- 115 (43) Institute for Land Water & Society, Charles Sturt University, Albury NSW 2640,  
116 Australia (masaunders@csu.edu.au)
- 117 (44) AgResearch, Lincoln Research Centre, Private Bag 4749, Christchurch 8140, New  
118 Zealand (nicole.schon@agresearch.co.nz)



(45) University of California Cooperative Extension, San Mateo & San Francisco  
Counties, Half Moon Bay, CA, USA (csidhu@alumni.ucsd.edu)

(46) Department of Zoology, Faculty of Science, Palacký University, Olomouc, Czech  
Republic (veselym@upol.cz)

(47) Terrestrial Ecology Research Group, Department for Ecology and Ecosystem  
Management, School of Life Sciences Weihenstephan, Technical University of Munich,  
Hans-Carl-von-Carlowitz-Platz 2, D-85354 Freising, Germany (wolfgang.weisser@tum.de)

**Corresponding author:** Elinor M. Lichtenberg; Department of Ecology & Evolutionary  
Biology, The University of Arizona, PO Box 210088, Tucson, AZ 85721, USA;  
elinor@email.arizona.edu; 1-240-393-0075

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DC, CMK, CK, and EML designed the study with support from FB, PB, RB, NAB-P, LGC,  
WES, NW, and RW; EML, DC, and CMK collected, prepared, and analyzed data and wrote  
the manuscript; all authors except CMK and EML contributed empirical field data; all  
authors revised the manuscript.



**ABSTRACT**

Agricultural intensification is a leading cause of global biodiversity loss, which can reduce the provisioning of ecosystem services in managed ecosystems. Organic farming and plant diversification are farm management schemes that may mitigate potential ecological harm by increasing species richness and boosting related ecosystem services to agroecosystems. What remains unclear is the extent to which farm management schemes affect biodiversity components other than species richness, and whether impacts differ across spatial scales and landscape contexts. Using a global meta-dataset, we quantified the effects of organic farming and plant diversification on abundance, local diversity (communities within fields), and regional diversity (communities across fields) of arthropod pollinators, predators, herbivores, and detritivores. Both organic farming and higher in-field plant diversity enhanced arthropod abundance, particularly for rare taxa. This resulted in increased richness but decreased evenness. While these responses were stronger at local relative to regional scales, richness and abundance increased at both scales, and richness on farms embedded in complex relative to simple landscapes. Overall, both organic farming and in-field plant diversification exerted the strongest effects on pollinators and predators, suggesting these management schemes can facilitate ecosystem service providers without augmenting herbivore (pest) populations. Our results suggest that organic farming and plant diversification promote diverse arthropod meta-communities that may provide temporal and spatial stability of ecosystem service provisioning. Conserving diverse plant and arthropod communities in farming systems therefore requires sustainable practices that operate both within fields and across landscapes.

## INTRODUCTION

Simplification of agricultural landscapes, and increased use of fertilizers and pesticides, threaten arthropod communities worldwide (Matson *et al.*, 1997; Tschamntke *et al.*, 2005; Potts *et al.*, 2016). This could impair agricultural sustainability because declines in arthropod abundance and diversity are often associated with reduced provisioning of ecosystem services including pollination, pest control, and nutrient cycling (Kremen & Miles, 2012; Oliver *et al.*, 2015). Two strategies purported to mitigate this ecological harm are organic farming and in-field plant diversification (Table S1). We refer to these strategies as farm management schemes, both of which include a host of practices that promote biological diversification (Kremen & Miles, 2012; Puech *et al.*, 2014). We refer to organic farming, conventional farming, high in-field plant diversification, and low in-field plant diversification as separate field types. Mounting evidence indicates that arthropod communities are more diverse and abundant in fields lacking synthetic fertilizers and pesticides, and in those with greater plant diversity (e.g., intercropped or having non-crop vegetation like hedgerows or floral strips) (Letourneau *et al.*, 2011; Crowder *et al.*, 2012; Kennedy *et al.*, 2013; Garibaldi *et al.*, 2014; Batáry *et al.*, 2015; Fahrig *et al.*, 2015).

The benefits of diversified farming practices may manifest at different scales, such as within individual fields (local diversity) or across multiple fields in a landscape (regional diversity) (Table S1). One observational study of 205 farms across Europe and Africa, for example, found that although organic farming provided strong benefits for local richness of plants and pollinators, these benefits faded at regional scales (Schneider *et al.*, 2014). This suggests that while farmers may promote local diversity on their field(s) by using organic practices, their efforts may not enhance biodiversity across multiple fields. Conversely, the

addition of hedgerows to crop fields has been shown to increase community heterogeneity and species turnover (measures of local diversity), which are important components of regional diversity (Ponisio *et al.*, 2016). The effects of farm management for particularly mobile arthropods, such as pollinators, may also transcend individual fields if the improved quality of habitats on one field boosts abundance, with organisms spilling over to nearby fields (Tscharrntke *et al.*, 2012; Kennedy *et al.*, 2013). While increases in local diversity have been shown to provide the strongest benefits to individual ecosystem services (i.e., pollination and biological control), regional diversity can support the simultaneous provision of multiple ecosystem services over space and time (Pasari *et al.*, 2013). Thus, to mitigate the effects of biodiversity loss across agroecosystems, farm management schemes should ideally benefit both local and regional diversity.

Research on the impacts of organic farming and in-field plant diversity has primarily focused on beneficial functional groups such as natural enemies and pollinators (Crowder *et al.*, 2010; Kennedy *et al.*, 2013) across intensively sampled regions of Europe and North America (Shackelford *et al.*, 2013; De Palma *et al.*, 2016). Moreover, almost all studies rely on richness (the number of taxa; Table S1) as a proxy for biodiversity but ignore metrics such as evenness (the relative abundances among species; Table S1) (e.g., Bengtsson *et al.*, 2005; Tuck *et al.*, 2014). Yet, richness poorly reflects overall community diversity (Duncan *et al.*, 2015; Loiseau & Gaertner, 2015), and its measurement is strongly confounded by abundance (Chao & Jost, 2012). Variation in richness has also been shown to have minimal impacts on ecosystem functioning when richness increases are driven primarily by rare species that contribute little to ecosystem services (Kleijn *et al.*, 2015; Winfree *et al.*, 2015). While common species may provide the majority of ecosystem services on some farms (Schwartz *et*

*al.*, 2000; Kleijn *et al.*, 2015), rare species can provide redundancy (Kleijn *et al.*, 2015) or support provisioning of multiple ecosystem services (Soliveres *et al.*, 2016). Assessing evenness can help determine whether richness increases are driven by rare or common species. Richness, evenness, and abundance can also independently or interactively affect ecosystem function (Wilsey & Stirling, 2006; Wittebolle *et al.*, 2009; Crowder *et al.*, 2010; Northfield *et al.*, 2010; Winfree *et al.*, 2015). Thus, teasing apart the effects of farm management schemes on abundance and each diversity metric is critical. While existing studies find that organic farming and in-field plant diversification tend to boost abundance and richness of certain taxa, whether these effects are consistent for other biodiversity components such as evenness, for functional groups other than pollinators and natural enemies, and for less-well studied regions of the world (e.g., the tropics and Mediterranean) remains unclear.

Here, we present a comprehensive synthesis of studies that explore how organic farming and in-field plant diversification influence arthropod communities across global agroecosystems. We determine whether community responses to these management schemes vary based on different metrics (abundance, local richness and evenness, regional richness and evenness) and arthropod functional groups (detritivores, herbivores, pollinators, and predators). We investigate if these responses depend on landscape complexity (i.e., the proportion of natural and semi-natural habitat surrounding the farm; Fig. S1, Table S1), because landscape heterogeneity has been shown to influence the effectiveness of farm management schemes (Batáry *et al.*, 2011; Kleijn *et al.*, 2011; Kennedy *et al.*, 2013; Tuck *et al.*, 2014). We also explore whether farm management schemes have similar impacts on relatively rare compared to common taxa. Our results demonstrate whether local and regional

diversity and abundance of different functional groups are similarly affected by on-farm management and landscape complexity, and the extent of covariance between biodiversity within and across fields in a landscape. Broadly, our findings further reveal the role of farm management in mitigating biodiversity loss and maintaining healthy arthropod communities in agroecosystems under global change.

## **MATERIALS AND METHODS**

### **Literature survey**

We compiled data from studies on arthropod diversity in agroecosystems that compared one or both of the farm management schemes of interest: (1) organic vs. conventional farming and (2) high vs. low in-field plant diversity. We defined organic agriculture as fields that were organically certified or met local certification guidelines (Table S1). These guidelines involve, at minimum maintaining production systems free of synthetic pesticides and fertilizers. We defined conventional agriculture as fields or farms that used recommended rates of synthetic, or a mix of synthetic and organic, pesticides and fertilizers. Other types of farming systems, such as integrated, which fit neither category were excluded from the analysis. Fields were defined as having high in-field plant diversity if they had diverse crop vegetation or managed field margins to include non-crop vegetation (e.g., hedgerows, border plantings, flower strips) (Table S1). We also classified small (< 4 ha) fields as diverse because they yield small-scale crop diversity (across several fields) even if the target field is a monoculture (Pasher *et al.*, 2013). Fields were defined as having low in-field plant diversity if they had none of these features. Studies that compared these schemes were identified by (1) searching the reference lists of recent meta-analyses (Batáry *et al.*,

2011; Chaplin-Kramer *et al.*, 2011; Crowder *et al.*, 2012; Garibaldi *et al.*, 2013; Kennedy *et al.*, 2013; Scheper *et al.*, 2013; Shackelford *et al.*, 2013), (2) searching ISI Web of Knowledge (April and May 2013) using the terms “evenness or richness” and “organic and conventional” or “local diversity”, and (3) directly contacting researchers who study arthropods in agricultural systems.

We identified 235 relevant studies that we examined for inclusion based on five criteria: (1) sampling was performed in the same crop or crop type (e.g., cereals) for organic and conventional fields, or fields with high and low in-field plant diversity; (2) sampling was conducted at the scale of individual crop fields rather than using plots on experiment stations; (3) the study included at least two fields of each type; (4) all organisms collected were identified to a particular taxonomic level (i.e., order, family, genus, species, or morphospecies), such that no taxa were lumped into groups such as “other”; and (5) at least three unique taxa were collected. We use “taxon” to refer to a single biological type (e.g., species, morphospecies, genus, family), determined as the finest taxonomic resolution to which each organism was identified in a particular study (see examples in Table S1). A total of 60 studies met our criteria, representing 43 crops, 21 countries, and 5 regions (Asia, Europe, North and Central America, South America, Oceania) (Fig. S2, Table S2). For studies that investigated both management scheme comparisons, we included the data in both analyses only when the field types were independently assigned (Table S3); otherwise we selected the scheme that the authors indicated the study was designed to address (Table S2). Across these 60 studies, our meta-analysis included 110 unique data points: 81 comparing organic and conventional fields and 29 comparing fields with high vs. low in-field plant diversity (Fig. S2, Tables S2, S4, archived data). Among organic vs. conventional studies, the

number with high in-field plant diversity, low in-field plant diversity, and both levels of plant diversity was independent of organic vs. conventional management ( $\chi^2_2 = 0.47$ ,  $p = 0.79$ ).

### Calculation of effect sizes

Unlike traditional meta-analyses that extract summary statistics from studies, we gathered and manipulated raw data, which enabled us to calculate evenness and classify taxa into functional groups. For each study, we compiled data on the abundance of all taxa in each field. For studies conducted across multiple years or crop types, separate values were compiled for each year and crop. To avoid pseudoreplication, for multi-year studies we selected a single year to analyze based on maximizing the number of (1) sites that met the evenness criterion (at least three taxa), (2) fields, or (3) individuals (in decreasing priority order; Garibaldi *et al.*, 2013). Each collected taxon was classified into one of four functional groups: detritivore, herbivore, pollinator, or predator (see Supporting Methods for details). These taxon-level data were used to calculate effect sizes for abundance, local diversity, and regional diversity in paired organic vs. conventional or high vs. low in-field plant diversity systems. For local and regional calculations, we defined diversity as both richness and evenness, and treated each functional group separately (Fig. S1).

Local diversity reflects the average diversity within each field, and was calculated using individual crop fields as the sampling unit (Fig. S1, Table S1). In studies with subsamples at a scale smaller than a field (i.e., plots within fields), values across these subsamples were averaged before calculating local diversity. Abundance was the number of arthropods, and richness the number of unique taxa, in a field. Evenness was calculated using the metric  $E_{var}$ , which ranges from 0 (one taxon dominant) to 1 (uniform abundance for all



301 taxa). This metric was chosen for its desirable statistical properties, particularly independence  
302 from richness, and its use in similar previous meta-analyses (Crowder *et al.*, 2012). After  
303 calculating abundance, richness, and evenness for each field, we averaged values across all  
304 fields of a particular type in a study to obtain the values for effect size calculations.

305 Regional diversity values were calculated based on individuals pooled across all fields  
306 in a study (Fig. S1, Table S1). Thus, regional richness and evenness are measures of diversity  
307 of meta-communities across fields in a landscape, while local diversity measures  
308 communities in a single field (Wang & Loreau, 2014). We note that regional diversity is not a  
309 direct indication of spatial scale, as the geographical extent of sampling varied among  
310 studies. Some studies were not designed to assess regional diversity specifically, and sampled  
311 unequal numbers of fields of each type. To correct for this sampling bias, we used sample-  
312 based rarefaction with 1,000 random samples taken from the set of fields in a given study to  
313 determine pooled species assemblages (Gotelli & Colwell, 2011). For example, if a study had  
314 10 conventional and 6 organic fields, regional diversity values for the conventional  
315 management schemes would be based on the average pooled community taken from 1,000  
316 random draws of 6 field sites. Regional abundance is simply local abundance multiplied by  
317 the number of sites, thus we reported only one abundance value per study.

318 To compare effects of farm management schemes on diversity and abundance, we  
319 used the log-response ratio as an effect size metric (Hedges *et al.*, 1999). We used this metric,  
320 rather than a weighted effect size, for three reasons. First, weighted effect sizes could not be  
321 calculated for regional diversity because these calculations were based on a single value  
322 (without replication) from each study, such that there was no estimate of variability. Second,  
323 our studies classified arthropods at varying levels of taxonomic resolution. Studies classified

at the family level had less variability than studies classified at the species level, so using a weighted metric would give studies conducted at a coarser taxonomic resolution greater weight. Finally, preliminary analysis showed weighted and unweighted analyses of local diversity and abundance were qualitatively similar (Table S5). In the Results, we back-transformed log response-ratio effect sizes to percentages.

We assessed funnel plot asymmetry to test for publication bias. Because we used an unweighted effect size metric, we plotted effect sizes against sample sizes (i.e., number of fields; Figs. S3, S4) (Sterne & Egger, 2001), and visually assessed asymmetry since formal statistical tests require effect size variances (Jin *et al.*, 2015) and measures of regional diversity had no variance component. Visual assessment looked for, and did not find, areas of missing non-significant results, a directional bias to effects, or a strong relationship between effect and sample sizes. We did not detect any sign of publication bias; funnel plots were sufficiently symmetrical. Finally, we ensured the sampling method (active versus passive sampling techniques) did not influence results (see Supporting Information, Table S6). We calculated abundance and diversity values with R v. 3.1.1 (R Core Team, 2014), using packages BiodiversityR (Kindt & Coe, 2005), doBy (Højsgaard & Halekoh, 2013), and reshape (Wickham, 2007).

## Study variables

We gathered data on three categorical variables and assessed whether they mediated arthropod responses to farm management schemes: (1) landscape complexity (simple, complex), (2) biome (boreal, Mediterranean, temperate, tropical), and (3) crop cultivation period (annual, perennial). Landscape complexity (see Fig. S1, Table S1) was determined

from land cover data on the percentage of natural and semi-natural habitat within 1 km of sampled fields. Natural and semi-natural habitat was defined as areas dominated by forest, grassland, shrubland, wetlands, ruderal vegetation, or non-agricultural plantings (i.e., previously-cultivated areas where vegetation is regenerating, hedgerows, field margins, and vegetation along roadways or ditches). For each study, we calculated the mean percentage of natural habitats across fields using locally-relevant land cover databases. Landscapes were classified as simple if they averaged  $\leq 20\%$  natural habitat, and complex if they averaged  $> 20\%$  natural habitat, following Tschardt et al. (2005) and common practice (e.g., Batáry *et al.*, 2011; Scheper *et al.*, 2013) (see Supporting Methods for additional details). Biome was based on the geographic location of the study. Crop cultivation periods were derived from several sources (FAO AGPC, 2000; Garibaldi *et al.*, 2013). Table S4 shows the distribution of data points across each of these descriptive variables.

## Data analyses

Table S7 summarizes specific questions we addressed and the approach we used to test each one. We first used one-sample *t*-tests (Crowder & Reganold, 2015) to determine if the mean effect sizes for abundance, local richness and evenness, and regional richness and evenness differed significantly from 0. For each management scheme comparison (organic vs. conventional or high vs. low in-field plant diversity), these analyses were conducted for the overall arthropod community and for each functional group separately. We also explored correlations between local and regional richness, and between local and regional evenness, to determine if these metrics responded similarly to each of the management schemes. We used  $\alpha = 0.10$ , to describe effect sizes that appeared ecologically important but did not meet the

somewhat arbitrary  $\alpha = 0.05$ . This accords with a recent policy statement by the American Statistical Association (Wasserstein & Lazar, 2016), which notes that reliance on arbitrary alpha values can lead to erroneous conclusions.

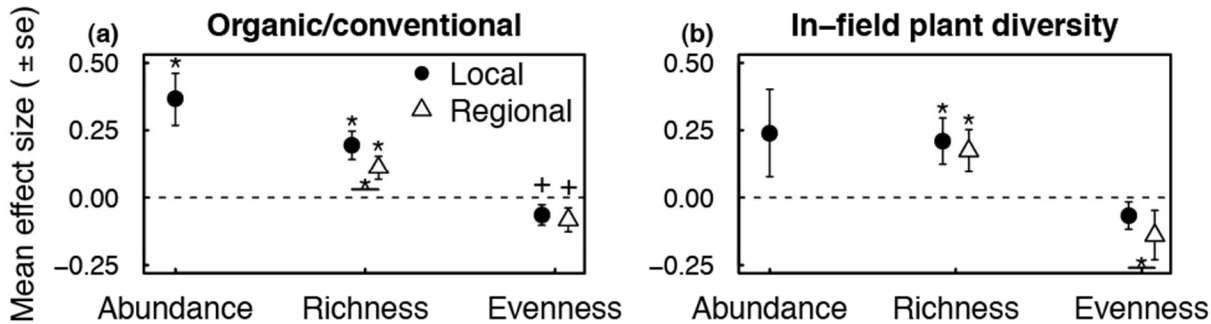
In subsequent analyses, we used meta-regression to examine whether effect sizes were influenced by functional group and other study characteristics. We excluded studies lacking landscape complexity data (see archived data) from meta-regressions. For each management scheme and response, we ran a linear mixed model (lme4 package; Bates *et al.*, 2014) that included eight fixed effect variables: (1) functional group (detritivore, herbivore, predator, pollinator), (2) diversity scale (local, regional), (3) landscape complexity (simple, complex), (4) biome (boreal, Mediterranean, temperate, tropical), (5) crop cultivation period (annual, perennial), (6) functional group×diversity scale interaction; (7) functional group×landscape complexity interaction; and (8) diversity scale×landscape complexity interaction. These models included study ID as a random effect. We used information-theoretic model selection to determine the set of best-fit models for each response variable (MuMIn package; Barton, 2014), which contained models with AICc values within 2 of the smallest value (Burnham & Anderson, 1998). We examined significance of the fixed effects in each model in the best-fit set ( $\alpha = 0.10$ ) with likelihood ratio tests, and used post-hoc planned contrasts (with *p*-values adjusted to control the overall Type I error rate using Holm's sequential Bonferroni procedure; see Supporting Methods) (phia package; Rosario-Martinez, 2013) to test for (1) differences in effect size among functional groups and biomes, (2) differences in effect size between the local and regional scales within each functional group, and (3) landscape complexity differences between each pair of functional groups.

We also tested whether abundance and richness effect sizes differed for rare and common taxa. Following Kleijn et al. (2015), within each study we classified taxa as common if their relative abundance was at least 5% of the total community; other species were categorized as rare. We then calculated local abundance and richness as well as regional abundance and richness separately for rare and common taxa. We used one-sample *t*-tests to determine if mean effect sizes differed significantly from zero, and paired *t*-tests to determine whether mean effect sizes differed between rare and common taxa.

## RESULTS

### Effects of management schemes on overall arthropod communities

Organic farming increased arthropod abundance (45% change), local richness (19%), and regional richness (11%) (Fig. 1a, Table S8). These positive effects were stronger for local compared to regional richness (Fig. 1a, Tables S9, S10). Arthropod communities on organic farms had significantly but only moderately lower local evenness (-6%) and regional evenness (-8%) than on conventional farms (Fig. 1a, Table S8). Fields with high in-field plant diversity increased local richness (23%) and regional richness (19%), with similar magnitude (Fig. 1b, Tables S8, S11, S12). In-field plant diversity did not significantly affect abundance (27%), local evenness (-6%) or regional evenness (-13%) (Fig. 1b, Table S8). Overall, there were strong positive correlations between local and regional richness ( $r = 0.87$ ), and between local and regional evenness ( $r = 0.57$ ; Fig. S5).



**Figure 1.** Effects of farm management schemes on arthropod abundance, local diversity, and regional diversity. Values shown are for the entire arthropod community, and represent the mean log-response ratio ( $\pm$  SE) of (a) adopting organic farming and (b) promoting in-field plant diversity on abundance, richness, and evenness. A “\*” ( $p < 0.05$ ) or “+” ( $0.05 \leq p < 0.1$ ) above a mean denotes a significant difference from zero (determined via one-sample  $t$ -tests; statistical details in Table S8), while one below a pair of means indicates a significant difference between local and regional diversity (determined via linear mixed models; Tables S9-S12).

Organic farming increased abundance and richness of both rare and common arthropods at the local and regional scales (Fig. S6a,c, Table S13). At the local scale, organic farming increased arthropod richness by promoting rare taxa (27% increase) more strongly than common taxa (14% increase) (Fig. S6c, Table S14). In-field plant diversification also had differential effects on rare and common taxa, increasing richness of both at the local scale, but only of rare taxa at the regional scale (Fig. S6d, Table S13). Fields with higher in-field plant diversity increased abundance of common arthropods, but not of rare arthropods (Fig. S6b, Table S13).

### Effects of management schemes on arthropod functional groups

Organic farming substantially increased the abundance (90%), local richness (55%), and regional richness (32%) of pollinator communities, but did not impact pollinator evenness (Fig. 2a, Table S15). For predator communities, organic farming increased abundance (38%) and local richness (14%), lowered local (-9%) and regional (-14%)

evenness (Fig. 2c, Table S16), but did not affect regional richness (Fig. 2c, Table S16). Organic farming also did not impact abundance, local or regional richness, or local or regional evenness for herbivore (Fig. 2e, Table S17) or detritivore (Fig. 2g, Table S18) communities. For all biodiversity components and functional groups, effect sizes in response to organic farming did not differ between the local and regional scales (Fig. 2a,c,e,f, Tables S9, S10). The diversity scale $\times$ landscape complexity interaction was never retained in a best-fit model (Tables S9, S11).

High in-field plant diversity promoted the abundance (45%), local richness (44%), and regional richness (29%) of pollinator communities, but decreased local pollinator evenness (-11%) (Fig. 2b, Table S15). In-field plant diversity did not affect regional pollinator evenness (Fig. 2b, Table S15). In addition, in-field plant diversity did not alter abundance, local or regional richness, or local or regional evenness for predator (Fig. 2d, Table S16) or herbivore (Fig. 2f, Table S17) communities. In-field plant diversity increased the regional richness (69%) of detritivores and lowered regional detritivore evenness (-65%), but did not impact detritivore abundance, local richness, or local evenness (Fig. 2h, Table S18). The low sample size for detritivores, however, limits our ability to make inferences about this group.

#### **Effects of landscape complexity, biome, and crop cultivation period on arthropod communities**

Landscape complexity did not mediate the influences of organic farming or in-field plant diversity on arthropod abundance or evenness (Fig. 3, Tables S9-S12). However, both management schemes had stronger positive effects on local and regional arthropod richness



460 in complex relative to simple landscapes: organic farming 26% vs. 9%, in-field plant  
461 diversification 29% vs. 11%, respectively (Fig. 3c,d, Tables S9-S12). The effects of  
462 landscape complexity were similar in both direction and magnitude for local and regional  
463 diversity (Fig. 3c-e, Tables S9-S12). Organic farming promoted herbivore richness to a  
464 greater extent in simple than complex landscapes (Table S10), but other effects of landscape  
465 complexity on abundance and diversity were similar across functional groups (Tables S9-  
466 S12).

467 Stronger richness gains in complex than simple landscapes were driven  
468 predominantly by rare taxa (Fig. 4). In complex landscapes, both organic farming and in-field  
469 plant diversification had stronger positive effects on local richness of rare (organic 44%,  
470 plant diversification 68%) than of common (organic 21%, plant diversification 18%)  
471 arthropod taxa (Fig. 4c,d, Table S19). Organic farming within complex landscapes also  
472 increased local abundance and regional richness of rare taxa (78% and 17%, respectively) to  
473 a greater extent than common taxa (33% and 4%, respectively) (Fig. 4a, Table S19). Neither  
474 management scheme differentially affected abundance or richness of rare and common taxa  
475 in simple landscapes (Fig. 4, Table S19).

476 Biome mediated the impacts of in-field plant diversity on arthropod richness (pooled  
477 across local and regional scales) (Tables S11, S12). Post-hoc tests failed to indicate  
478 significant differences among biomes when considering all studies; but when the single  
479 boreal study was removed from the analysis, high in-field plant diversity more strongly  
480 promoted richness in Mediterranean (53%) than in temperate studies (-2%) (Table S12).  
481 Biome did not mediate the effects of organic farming or in-field plant diversification on  
482 arthropod abundance or evenness (Tables S9-S12). Organic farming increased arthropod

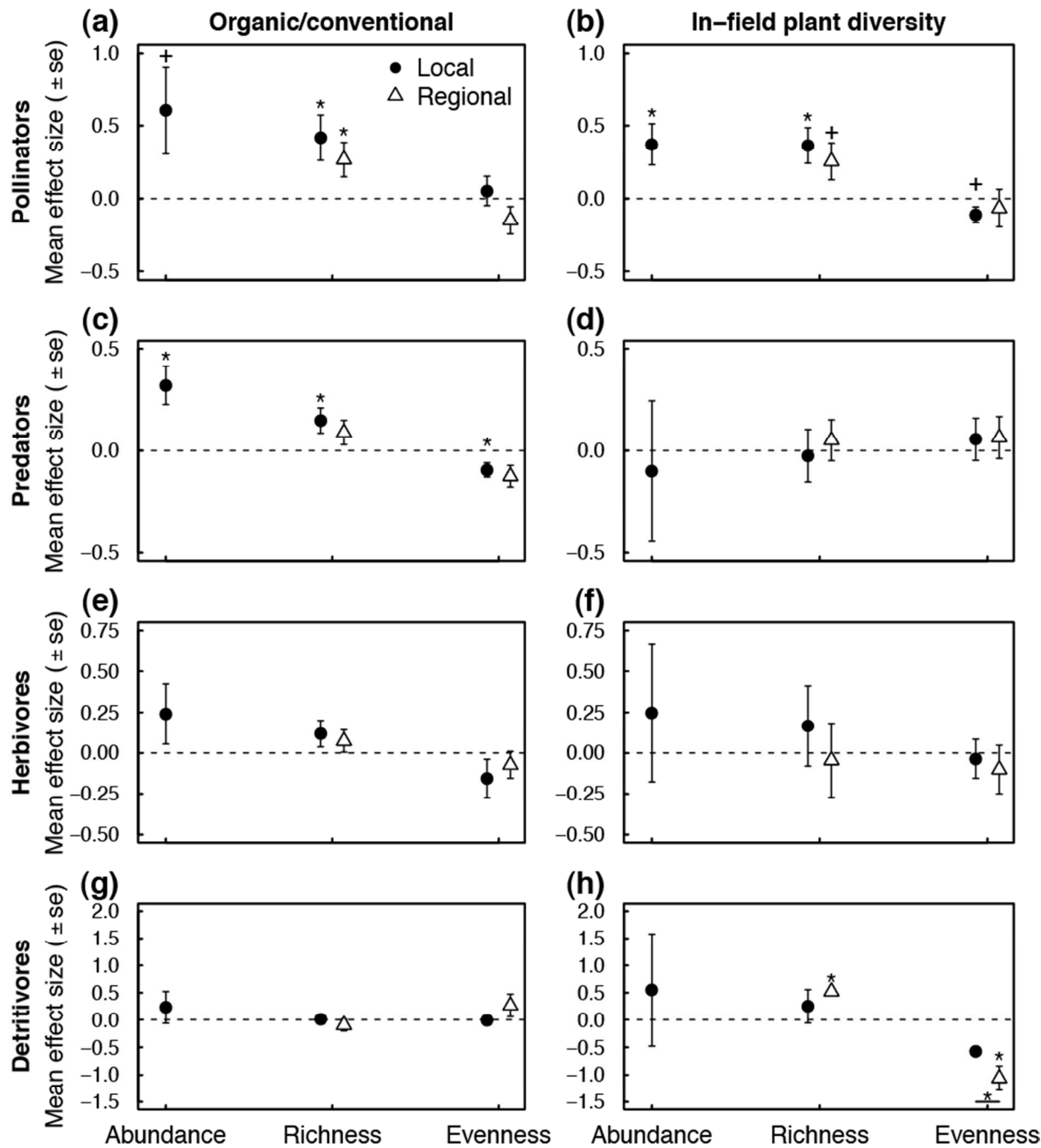
483 abundance to a greater extent in annual (70%) than in perennial (1%) crops (Tables S9, S10).

484 The effects of in-field plant diversification on abundance and diversity were consistent across

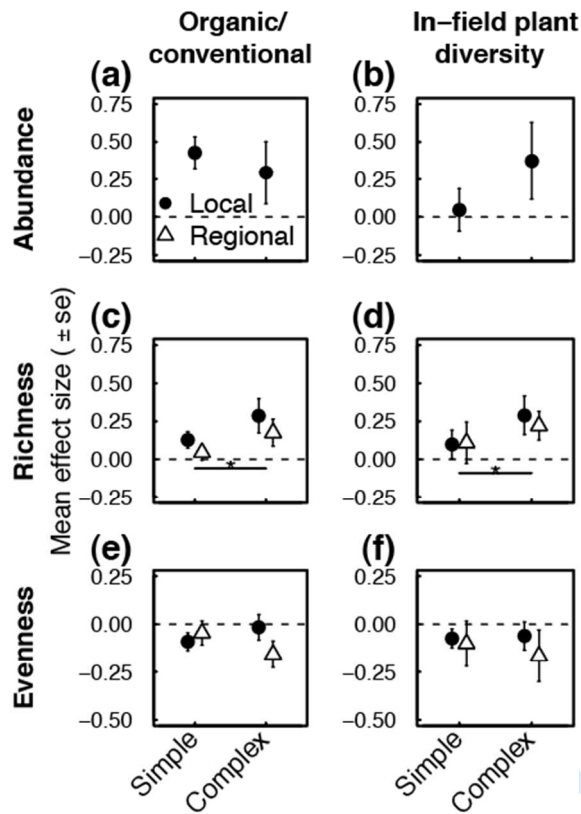
485 crop cultivation periods (Tables S11, S12).

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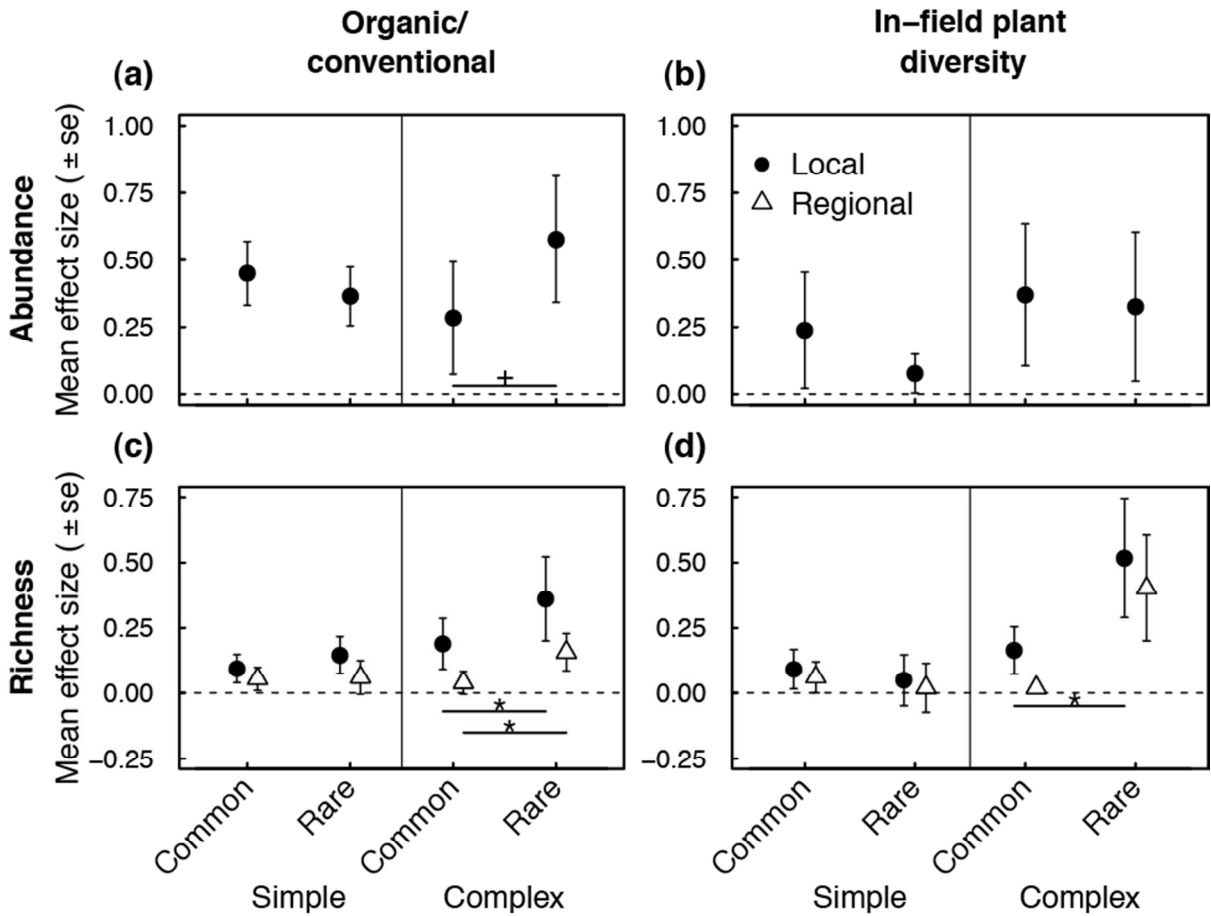
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**Figure 2.** Effects of farm management schemes on abundance, local diversity, and regional diversity of arthropod functional groups. Mean log-response ratios ( $\pm$  SE) of (left column) adopting organic farming and (right column) promoting in-field plant diversity for (a-b) pollinators, (c-d) predators, (e-f) herbivores, and (g-h) detritivores. A “\*” ( $p < 0.05$ ) or “+” ( $0.05 \leq p < 0.1$ ) above a mean denotes a significant difference from zero (determined via one-sample  $t$ -tests; Tables S15-S18). Meta-regressions indicated that differences between local and regional values did not vary with functional group (Tables S9-S12).



**Figure 3.** Effects of landscape complexity on the entire arthropod community in organic vs. conventional farms (left column) and fields with high vs. low in-field plant diversity (right column). Each graph shows the mean log-response ratio ( $\pm$  SE) for studies in simple ( $\leq 20\%$  natural habitat) or complex ( $>20\%$  natural habitat) landscapes for (a,b) abundance, (c,d) richness, and (e,f) evenness. A “\*” ( $p < 0.05$ ) or “+” ( $0.05 \leq p < 0.1$ ) below a set of means indicates a significant difference between means at the habitat complexity levels (Tables S9-S12).



**Figure 4.** Effects of farm management schemes on abundance (a, b) and richness (c, d) of common vs. rare taxa in simple and complex landscapes. Mean log-response ratios (±SE) of (left column) adopting organic farming and (right column) promoting in-field plant diversity. A “\*” ( $p < 0.05$ ) or “+” ( $0.05 \leq p < 0.1$ ) below a pair of means indicates a significant difference between rare and common taxa within a landscape complexity category (determined via paired  $t$ -tests; Table S19).

## DISCUSSION

Our global meta-analysis showed that both organic farming and in-field plant diversification strongly increased arthropod abundance and richness, but had weaker effects on evenness. The minimal evenness decreases on diversified farms reflected the presence of more rare taxa. Emerging evidence suggests that rare taxa contribute to individual ecosystem services less than common taxa (Schwartz *et al.*, 2000; Kleijn *et al.*, 2015), although they

may be important for maintenance of multiple ecosystem services across time and space (Isbell *et al.*, 2011; Soliveres *et al.*, 2016). Thus, while organic farming and plant diversification promote arthropod biodiversity conservation goals, their impacts on ecosystem services may be nuanced. The positive effects of both organic farming and in-field plant diversification were greatest for two groups of beneficial arthropods: pollinators and predators. Thus, both schemes may increase agroecosystem sustainability by promoting key ecosystem service providers without boosting pest (herbivore) densities.

Previous meta-analyses have investigated how organic farming and, to a lesser extent, in-field plant diversification, affect arthropod abundance and richness (e.g., Bengtsson *et al.*, 2005; Batáry *et al.*, 2011; Chaplin-Kramer *et al.*, 2011; Scheper *et al.*, 2013; Shackelford *et al.*, 2013; Tuck *et al.*, 2014). Our study extends upon this work by (1) combining data on multiple arthropod functional groups (but see Shackelford *et al.*, 2013), and (2) examining the type and scale of diversity across a variety of crop types. As such, we offer a more comprehensive understanding of when and how farm management schemes alter arthropod biodiversity. Our findings caution that the frequent use of richness as the sole proxy for biodiversity fails to reflect the full impacts of farming practices on biologic communities. While multiple studies have shown that organic farming boosts richness (e.g., Bengtsson *et al.*, 2005; Tuck *et al.*, 2014), we found that evenness decreased, an outcome that was due mainly to promotion of rare species. Species richness might be increased by conservation practices that target specific taxa, but the promotion of evenness requires practices that can simultaneously balance the abundances of many taxa (Crowder *et al.*, 2010, 2012). Finally, our results highlight the necessity of targeting farm management within the context of local conditions (Cunningham *et al.*, 2013; Saunders *et al.*, 2016). For example, our results suggest

that farmers in Mediterranean biomes might see greater arthropod richness gains by increasing in-field plant diversity than by farming organically, while farmers growing annual crops may be more likely to boost arthropod abundance with organic farming.

Disentangling relationships between biodiversity components at local and regional scales can inform patterns of community assembly and mechanisms that shape community structure (Gering & Crist, 2002; Wang & Loreau, 2014). We found that regional diversity positively correlated with local diversity under both management schemes. Further, organic farming increased richness at both scales, although local effects were stronger than regional ones. One possible explanation is that diversified farming practices increase the heterogeneity of local communities (e.g., Ponisio *et al.*, 2016), which could lead to greater regional diversity. Another possibility is that diversified fields serve as source habitats within a matrix of crop and non-crop habitats across farming landscapes (M'Gonigle *et al.*, 2015). Further, the benefits of diversification practices on local communities in fields can be strongly mediated by regional species pools across farming landscapes (Gering & Crist, 2002).

Our results, in combination with another recent meta-analysis (Schneider *et al.*, 2014), suggest that mobility of organisms can determine whether the benefits of farm diversification accrue at both local and regional scales. While we show that organic farming can boost arthropod diversity at local and regional scales, Schneider *et al.* (2014) found that organic farming increased plant, earthworm, and spider richness at field but not regional scales. These groups of organisms tend to have limited dispersal capacity, particularly plants and earthworms. Thus, their local communities may be structured more by competition than long-distance dispersal (Gering & Crist, 2002), which would limit the similarity between communities within and across fields. At the same time, Schneider *et al.* (2014) found that



organic farming boosted the richness of bees, a more mobile group of organisms, by approximately 25% at the local scale and 15% at the regional scale. We likewise found that diversified farming increased abundance, and local and regional richness, of mobile pollinators, but had less impact on detritivores that tend to have lower mobility (Sattler *et al.*, 2010).

Overall, our results are consistent with mounting evidence that farm management and landscape complexity interactively affect arthropod biodiversity (e.g., Rusch *et al.*, 2010; Batáry *et al.*, 2011; Kennedy *et al.*, 2013; Tuck *et al.*, 2014), although results across studies have found sometimes conflicting patterns (Kleijn *et al.*, 2011; Tscharntke *et al.*, 2012; Tuck *et al.*, 2014). For example, agri-environment schemes that promote low input, low disturbance, and diverse farms are sometimes most effective in fostering biodiversity in structurally simple landscapes (Batáry *et al.*, 2011; Scheper *et al.*, 2013). This presumably occurs because simple landscapes fail to satisfy the resource needs of many species, such that these species may disperse into diverse farms to seek resources (Tscharntke *et al.*, 2005; Kremen & Miles, 2012). In contrast, we found that impacts of organic farming and plant diversification on arthropod richness were heightened for fields embedded in complex landscapes. This could occur if complex landscapes support more diverse species pools that can respond positively to farm management (Duelli & Obrist, 2003; Hillebrand *et al.*, 2008; Kennedy *et al.*, 2013). Consistent with this hypothesis, we showed that organic farming in complex landscapes preferentially increased richness of rare taxa locally (i.e., in fields) and regionally (i.e., across landscapes). Importantly, the interactive effects of landscape complexity and on-farm management may differ across arthropod functional groups with varying capacity to move across landscapes (Tscharntke *et al.*, 2005; Chaplin-Kramer *et al.*,

2011). However, the only interaction between landscape complexity and management schemes we found was for richness of herbivores, a group with considerable variation in mobility among taxa (Sattler *et al.*, 2010).

Ideally, increases in abundance and diversity of arthropods on farms would enhance the provisioning of ecosystem services (Kremen & Miles, 2012). However, empirical studies have provided mixed evidence. In-field plant diversification and increased landscape complexity have been found to promote predator abundance and diversity with no change in pest control levels (Chaplin-Kramer *et al.*, 2011; Rusch *et al.*, 2016) or reduced crop damage (Letourneau *et al.*, 2011). The relationship between biodiversity and ecosystem services on farms is thus likely strongly mediated by species' abundances and functional roles. For example, Northfield *et al.* (2010) found that greater predator richness increased pest control, but only with high predator densities where complementarity among predator species was fully realized. Increases in pollinator richness can have minimal impacts on ecosystem services when richness gains are associated with rare species that contribute little to pollination (Kleijn *et al.*, 2015; Winfree *et al.*, 2015). Increasing wild pollinator richness on large farms (> 14 ha) only increases fruit set when wild pollinator density is also high (Garibaldi *et al.*, 2016). Higher predator species evenness on organic farms has also been shown to translate to increased pest control, with the potential to reduce yield gaps compared with conventional agriculture (Crowder *et al.*, 2010). However, models suggest that decreased evenness could also lead to greater ecosystem services when abundance of common species that are effective ecosystem services providers increases at the expense of rare species that are functionally less important (Crowder & Jabbour, 2014), a result seen with pollinators in agricultural systems (Kleijn *et al.*, 2015; Winfree *et al.*, 2015). The

combination of context-specific responses to farm management schemes shown by this study and biodiversity-ecosystem functioning relationships that depend on species' abundances and functional traits suggest that the effects of diversified farming on ecosystem services are likely to depend on biome, landscape, and crop characteristics.

By promoting biodiversity and abundance of arthropods, diversified agriculture could provide a multitude of other benefits (Oliver *et al.*, 2015). Biodiversity can help maintain stability of ecosystem processes through mechanisms such as response diversity and functional redundancy (Cardinale *et al.*, 2012; Mori *et al.*, 2013). Arthropod richness gains in response to organic farming and plant diversification, such as those documented here, could guard against the loss of ecological function by supporting multiple species that occupy similar functional niches (functional redundancy) or that are functionally similar but respond differentially to environmental change (response diversity; Elmqvist *et al.*, 2003). The abundance and richness increases we detected for pollinators and predators but not for herbivores suggest that the two former groups may benefit more from these stabilizing processes. Resilient systems must also exhibit multiple ecosystem functions (multifunctionality) as environmental conditions and arthropod populations fluctuate. Increases in rare taxa, as detected in this study, may be critical for multifunctionality (Isbell *et al.*, 2011; Soliveres *et al.*, 2016) and even for single ecosystem functions (Zavaleta & Hulvey, 2004; Mouillot *et al.*, 2013). Thus, regional-scale refuges for rare species may ensure resilient agricultural systems.

Overall, our results suggest that organic farming and in-field plant diversification both promote biodiversity on farms. Moreover, these two schemes might have interactive effects on farm productivity. Practices such as multi-cropping (plant diversification) and longer,

more diverse, crop rotations can reduce the yield gaps between organic and conventional agriculture (Ponisio *et al.*, 2015), and increase the profitability of organic relative to conventional systems (Crowder & Reganold, 2015). Diversified small farms are increasingly being replaced by large, simplified, and intensive monoculture production systems (Tscharntke *et al.*, 2005; Bennett *et al.*, 2012). This is problematic because intensified farming reduces the long-term sustainability of agroecosystems, thereby threatening global food security (Ray *et al.*, 2012). One of the greatest challenges of the 21<sup>st</sup> century is meeting the food, fiber, and energy needs of a growing human population while maintaining farm sustainability and ecosystem functioning (Tilman *et al.*, 2011). Our study underscores that adopting organic farming or in-field plant diversification practices might aid society in attaining these goals.

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**DATA ACCESSIBILITY**

Data and scripts available at: [insert DOI for Zenodo repository]

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**SUPPORTING METHODS**

**Functional group classifications**

Data providers determined the functional group of each taxon. When functional groups were non-defined or non-standard (e.g., saprophage), or when taxa filled multiple functional roles (e.g., species that serve as both pollinators and herbivores), we assigned taxa to a single functional role based on their most common description in the literature. To maximize data inclusion, we also (1) combined predators and parasitoids, (2) classified all carabids as predators since even the herbivorous species are thought to consume some animal material (e.g., Hengeveld, 1980; Jørgensen & Toft, 1997), and (3) classified a few pollinators as herbivores in studies with few pollinator taxa but many herbivores.

**Sampling methods**

Studies used a broad range of sampling methods, which we categorized as active or passive. Active sampling methods included beating, netting bees seen at plants, hand-collecting individuals off plants, observational counting, washing plants, taking soil cores, sweep-netting, and vacuum sampling. Passive sampling methods were blue vane traps, light traps, visually-attractive or scented lures, malaise traps, minnow traps, pan traps, pitfall traps, and sticky cards. However, we did not include sampling method in our meta-regressions because preliminary analyses indicated that sampling method negligibly affected effect sizes (Table S5).

**Landscape complexity**

The “simple” landscape complexity category combined Tscharnkte et al.’s (2005) “cleared” and “simple” categories because we had only two “cleared” studies. We were unable to categorize landscape complexity when we obtained data directly from published articles that lacked GPS coordinates of sampling locations or information on natural habitat surrounding fields (Study IDs drit01, febe01, hesl01, hokk01, and weib01). These five studies all compared organic and conventional farms. In a couple of cases we based landscape complexity on percentage of natural habitat within 500 m (bosq01), or the average of percentages at 500m and 1.5 km (leto01; percentages at the two distances strongly correlated, with  $r = 0.8$ ).

**SUPPORTING REFERENCES**

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**Table S2 is in a separate file.**

**Table 1:** Definitions and descriptions of key terms.

Term	Definition	Notes
Organic farming	Organically certified, or meeting local certification guidelines. While guidelines vary by country, they typically involve, at minimum, maintaining production systems free of synthetic pesticides and fertilizers.	Both organic and conventional farming include a broad range of management strategies and levels of intensity (e.g., pesticide application frequency, monoculture vs. polyculture) (Kremen & Miles, 2012; Puech <i>et al.</i> , 2014).
Conventional farming	Fields or farms that used recommended rates of synthetic, or synthetic and organic, pesticides and fertilizers.	
In-field plant diversification	This includes various schemes that increase small-scale plant diversity, including intercropping, managing field margins to include non-crop vegetation (e.g., hedgerows, border plantings, flower strips), and use of small (< 4 ha) fields.	
Taxon	A single biological type (e.g., species, morphospecies, genus, family), determined as the finest taxonomic resolution to which each organism was identified.	Examples: <i>Apis mellifera</i> (species), <i>Halictus</i> sp. 1 (morphospecies), <i>Lasioglossum</i> spp. (genus), Formicidae (family). We assigned each taxon to a functional group (detritivore, herbivore, pollinator, predator), but calculated abundance and diversity from taxon-level data.
Abundance	The number of total individuals, of all taxa together, sampled.	We calculated abundance, richness, and evenness separately for each field type (e.g., conventional farming), crop, year, and arthropod functional group within each study.
Richness	The number of taxa sampled.	
Evenness	How individuals are distributed across taxa in the sample. The evenness measure that we used, $E_{var}$ , range from 0 (completely uneven, one taxon dominates) to 1 (completely even, with each taxon represented by an equal number of individuals).	
Region	A large spatial extent that contains multiple communities and habitats. We defined each study's region as	

	all of the fields sampled in the study.	
Rare taxon	A taxon with relative abundance less than 5% of all individuals sampled across the region.	We determined rarity separately for each management scheme comparison (organic vs. conventional, high vs. low in-field plant diversity), crop, year, and function within a study, but did not further separate by field type.
Local diversity	Diversity (here, richness and evenness) of a community within a field.	We estimated local abundance and diversity by first calculating abundance and diversity values within each field, then averaging these values across fields. For example, assume species A, B, C, D, and E were found in field 1; species A, E, and F in field 2; species B, C, D, and E in field 3; and species A, B, E, F, G, and H in field 4. Each field's richness would be 5, 3, 4, and 6, respectively. Local richness would be 4.5, the average of each field's richness value.
Regional diversity	Diversity (here, richness and evenness) of the meta-community that spans all fields in a region.	We estimated regional diversity by pooling individuals sampled in all fields within a landscape, then calculating diversity of taxa in this one regional sample. In the above example, the regional species pool would include species A through H and regional richness would be 8.
Landscape complexity	The proportion of natural and semi-natural habitat (areas dominated by forest, grassland, shrubland, wetlands, ruderal vegetation, or non-agricultural plantings including previously-cultivated areas where vegetation is regenerating, hedgerows, field margins, and vegetation along roadways or ditches) surrounding a farm.	We determined landscape complexity separately for each management scheme comparison, crop, and year within a study, by averaging proportions across fields.

**Table S3.** Fisher exact tests for studies with variation in both management (organic vs. conventional) and in-field plant diversity (high vs. low). These tests were used to determine whether sites were assigned independently to management types across the two management schemes. I-f=in-field plant diversity

Study ID	Number of sites with:				<i>p</i> -value	Management scheme(s) used
	Organic & high i-f	Organic & low i-f	Conventional & high i-f	Conventional & low i-f		
bomm01	8	16	22	53	0.80	Both
bosq01	7	10	10	10	0.74	Both
clou01	15	6	10	11	0.21	Both
danf01	2	0	3	5	0.44	Both
eige01	3	0	0	3	0.10	Organic/ conventional
ekro01	7	8	12	4	0.15	Both
frei01	0	2	2	0	0.33	I-f
frei02	2	0	0	2	0.33	I-f
holz01	16	5	10	11	0.11	Both
krem01	8	1	8	12	<0.0001	Organic/ conventional
leto01	5	0	0	5	0.0080	Organic/ conventional
ober01	3	2	0	3	0.20	Both
otie01	4	1	5	2	1.00	Both
rose01	0	12	9	0	<0.0001	Organic/ conventional
saun01	5	0	0	10	0.0003	I-f
weis01	1	6	3	22	1.00	Both

**Table S4.** Number of data points grouped by several categories used in the analysis.

<b>(a) Arthropod functional group</b>				
<b>Management scheme</b>	<b>Detritivore</b>	<b>Herbivore</b>	<b>Pollinator</b>	<b>Predator</b>
Organic/conventional	8	17	20	36
In-field plant diversity	3	5	13	8
<b>(b) Landscape complexity</b>				
<b>Management scheme</b>	<b>Simple</b>	<b>Complex</b>	<b>No data</b>	
Organic/conventional	44	30	7	
In-field plant diversity	12	17	0	
<b>(c) Biome</b>				
<b>Management scheme</b>	<b>Boreal</b>	<b>Mediterranean</b>	<b>Temperate</b>	<b>Tropical</b>
Organic/conventional	2	14	58	7
In-field plant diversity	1	9	13	6
<b>(d) Cultivation period</b>				
<b>Management scheme</b>	<b>Annual</b>	<b>Perennial</b>		
Organic/conventional	59	22		
In-field plant diversity	20	9		

**Table S5.** Correlations between unweighted (log-response ratio) and weighted (Hedges' *d*) effect sizes with various metrics. Weighted metrics could not be calculated at the regional scale (see Methods in main text)

Management scheme	Metric	Pearson's correlation coefficient	<i>t</i>	df	<i>p</i> -value
Organic vs. conventional	Abundance	0.66	7.88	79	<0.0001
Organic vs. conventional	Local richness	0.77	10.7	77	<0.0001
Organic vs. conventional	Local evenness	0.70	7.99	66	<0.0001
In-field plant diversity	Abundance	0.90	10.7	27	<0.0001
In-field plant diversity	Local richness	0.81	7.26	27	<0.0001
In-field plant diversity	Local evenness	0.83	7.01	22	<0.0001



66 **Table S6.** Effects of sampling method on effect size (log-response ratio) estimates. ANOVAs testing whether sampling method  
67 affected effect sizes were significant in only 4% of cases, which is within the amount expected by chance. Means are average  
68 untransformed log-response ratios comparing organic to conventional, or high to low in-field plant diversity, data. Effect sizes  
69 transformed to percentage change are in parentheses.  
70

Organic vs. conventional												
Functional group	Metric	N active	N passive	N both	Mean active	Mean passive	Mean both	SE active	SE passive	SE both	F	p-value
All	Abundance	32	39	10	0.56 (75%)	0.19 (21%)	0.43 (54%)	0.22	0.08	0.14	1.63	0.20
All	Local richness	32	39	10	0.29 (33%)	0.14 (15%)	0.11 (12%)	0.12	0.04	0.07	1.06	0.35
All	Local evenness	28	35	10	-0.07 (-6%)	-0.04 (-4%)	-0.12 (-12%)	0.07	0.05	0.05	0.22	0.80
All	Regional richness	32	39	10	0.16 (17%)	0.08 (9%)	0.07 (7%)	0.09	0.05	0.07	0.41	0.66
All	Regional evenness	32	39	10	-0.22 (-20%)	-0.00 (0%)	0.04 (5%)	0.08	0.05	0.06	3.62	0.031
Detritivore	Abundance	3	2	3	-0.37 (-31%)	0.83 (130%)	0.43 (54%)	0.51	0.63	0.29	1.70	0.27
Detritivore	Local richness	3	2	3	0.00 (0%)	0.17 (19%)	-0.09 (-9%)	0.14	0.17	0.07	1.00	0.43
Detritivore	Local evenness	3	1	3	0.20 (23%)	-0.04 (-4%)	-0.21 (-19%)	0.13	NA	0.07	4.15	0.11
Detritivore	Regional richness	3	2	3	0.00 (0%)	-0.38 (-32%)	0.00 (0%)	0.11	0.31	0.16	1.26	0.36
Detritivore	Regional evenness	3	2	3	0.22 (24%)	0.66 (94%)	0.05 (5%)	0.11	0.96	0.08	0.60	0.58
Herbivore	Abundance	8	6	3	0.29 (34%)	0.09 (10%)	0.39 (47%)	0.36	0.20	0.28	0.17	0.85
Herbivore	Local richness	8	6	3	0.04 (4%)	0.16 (17%)	0.24 (27%)	0.16	0.09	0.05	0.41	0.67

Herbivore	Local evenness	7	4	3	-0.23 (-20%)	-0.17 (-15%)	0.03 (3%)	0.21	0.19	0.09	0.32	0.73
Herbivore	Regional richness	8	6	3	-0.05 (-5%)	0.18 (20%)	0.19 (21%)	0.11	0.10	0.05	1.53	0.25
Herbivore	Regional evenness	8	6	3	-0.25 (-22%)	0.06 (6%)	0.12 (13%)	0.14	0.07	0.13	2.42	0.13
Pollinator	Abundance	12	7	1	0.98 (166%)	-0.10 (-9%)	1.06 (187%)	0.45	0.26	NA	1.56	0.24
Pollinator	Local richness	12	7	1	0.50 (64%)	0.28 (32%)	0.41 (51%)	0.25	0.15	NA	0.20	0.82
Pollinator	Local evenness	10	6	1	0.02 (2%)	0.18 (20%)	-0.39 (- 33%)	0.09	0.24	NA	0.90	0.43
Pollinator	Regional richness	12	7	1	0.26 (30%)	0.25 (29%)	0.36 (44%)	0.19	0.11	NA	0.02	0.98
Pollinator	Regional evenness	12	7	1	-0.16 (- 15%)	-0.11 (-10%)	-0.25 (- 22%)	0.15	0.11	NA	0.06	0.94
Predator	Abundance	9	24	3	0.54 (72%)	0.24 (28%)	0.27 (31%)	0.30	0.08	0.20	0.95	0.40
Predator	Local richness	9	24	3	0.32 (38%)	0.09 (10%)	0.08 (8%)	0.20	0.05	0.12	1.35	0.27
Predator	Local evenness	8	24	3	-0.13 (- 12%)	-0.08 (-8%)	-0.10 (- 9%)	0.11	0.04	0.06	0.17	0.85
Predator	Regional richness	9	24	3	0.26 (30%)	0.04 (5%)	-0.07 (- 7%)	0.17	0.06	0.09	1.63	0.21
Predator	Regional evenness	9	24	3	-0.42 (- 34%)	-0.04 (-4%)	0.06 (7%)	0.15	0.04	0.10	7.14	0.003
<b>In-field plant diversity</b>												
<b>Functional group</b>	<b>Metric</b>	<b>N active</b>	<b>N passive</b>	<b>N both</b>	<b>Mean active</b>	<b>Mean passive</b>	<b>Mean both</b>	<b>SE active</b>	<b>SE passive</b>	<b>SE both</b>	<b>F</b>	<b>p-value</b>
All	Abundance	13	11	5	0.22 (25%)	0.20 (22%)	0.37 (45%)	0.15	0.39	0.24	0.07	0.94
All	Local richness	13	11	5	0.29 (34%)	0.09 (10%)	0.26 (30%)	0.13	0.16	0.06	0.60	0.56

All	Local evenness	12	9	5	-0.03 (-3%)	-0.09 (-9%)	-0.11 (-10%)	0.07	0.11	0.05	0.25	0.78
All	Regional richness	13	11	5	0.25 (28%)	0.08 (9%)	0.19 (20%)	0.13	0.14	0.09	0.44	0.65
All	Regional evenness	13	11	5	-0.08 (-8%)	-0.25 (-22%)	-0.04 (-4%)	0.13	0.17	0.16	0.44	0.65
Detritivore	Abundance	1	2	0	0.03 (3%)	0.81 (125%)	NA	NA	1.73	NA	0.07	0.84
Detritivore	Local richness	1	2	0	-0.07 (-7%)	0.41 (51%)	NA	NA	0.45	NA	0.39	0.65
Detritivore	Local evenness	0	1	0	NA	-0.57 (-44%)	NA	NA	NA	NA	NA	NA
Detritivore	Regional richness	1	2	0	0.41 (50%)	0.58 (79%)	NA	NA	0.06	NA	2.55	0.36
Detritivore	Regional evenness	1	2	0	-0.84 (-57%)	-1.17 (-69%)	NA	NA	0.33	NA	0.32	0.67
Herbivore	Abundance	1	3	1	-0.04 (-4%)	0.30 (35%)	0.37 (45%)	NA	0.76	NA	0.03	0.97
Herbivore	Local richness	1	3	1	0.12 (13%)	0.15 (17%)	0.24 (27%)	NA	0.45	NA	0.01	0.99
Herbivore	Local evenness	1	2	1	0.21 (23%)	-0.17 (-16%)	0.00 (0%)	NA	0.19	NA	0.71	0.64
Herbivore	Regional richness	1	3	1	-0.06 (-6%)	-0.10 (-10%)	0.15 (16%)	NA	0.40	NA	0.05	0.95
Herbivore	Regional evenness	1	3	1	0.09 (10%)	-0.25 (-22%)	0.15 (17%)	NA	0.21	NA	0.61	0.62
Pollinator	Abundance	10	0	3	0.37 (46%)	NA	0.36 (43%)	0.15	NA	0.44	0.00	0.96
Pollinator	Local richness	10	0	3	0.40 (49%)	NA	0.26 (29%)	0.16	NA	0.12	0.22	0.65
Pollinator	Local evenness	10	0	3	-0.09 (-9%)	NA	-0.17 (-16%)	0.07	NA	0.07	0.39	0.55

Pollinator	Regional richness	10	0	3	0.28 (32%)	NA	0.17 (18%)	0.16	NA	0.16	0.13	0.73
Pollinator	Regional evenness	10	0	3	-0.05 (-5%)	NA	-0.10 (-10%)	0.15	NA	0.28	0.03	0.88
Predator	Abundance	1	6	1	-0.87 (-58%)	-0.05 (-5%)	0.40 (50%)	NA	0.44	NA	0.37	0.71
Predator	Local richness	1	6	1	-0.22 (-20%)	-0.05 (-5%)	0.30 (36%)	NA	0.16	NA	0.47	0.65
Predator	Local evenness	1	6	1	0.40 (49%)	0.01 (1%)	-0.03 (-3%)	NA	0.13	NA	0.71	0.54
Predator	Regional richness	1	6	1	0.08 (8%)	0.01 (1%)	0.27 (31%)	NA	0.13	NA	0.30	0.75
Predator	Regional evenness	1	6	1	0.23 (25%)	0.06 (6%)	-0.04 (-4%)	NA	0.14	NA	0.17	0.85

**Table S7:** Questions investigated in this study, and statistical tests that addressed each one. Q2, Q4, Q7, and Q8 were tested with the same meta-regression.

Question	How tested
(Q1) Does diversified farming differentially alter abundance, richness, and evenness?	One sample <i>t</i> -tests: Does each metric's mean effect size differ from zero?
(Q2) Diversified farming differentially alters local and regional diversity (richness, evenness).	(a) One-sample <i>t</i> -tests: Are patterns of difference from zero the same at the local and regional scales? (b) Meta-regression: Does scale affect mean effect size?
(Q3) Diversified farming differentially alters abundance and diversity of arthropods in different functional groups	One-sample <i>t</i> -tests: Within each functional group (detritivores, herbivores, pollinators, predators), does each metric's mean effect size differ from zero?
(Q4) Landscape complexity mediates responses of arthropod communities to diversified farming.	Meta-regression: Do effect sizes differ in simple and complex landscapes?
(Q5) Diversified farming differentially affects the abundance and diversity of relatively rare and relatively common taxa.	(a) One-sample <i>t</i> -tests: Does each metric's mean effect size for a given rarity category (rare, common) differ from zero? (b) Paired <i>t</i> -tests: Within a metric, do mean effect sizes for rare taxa differ from those of common taxa?
(Q6) Landscape complexity mediates the degree to which diversified farming differentially affects the abundance and diversity of rare vs. common taxa.	Paired <i>t</i> -tests: Within each metric and landscape complexity category (simple, complex), do mean effect sizes for rare taxa differ from those of common taxa?
(Q7) A crop's cultivation period (annual, perennial) mediates responses of arthropod communities to diversified farming.	Meta-regression: Do effect sizes differ for crops grown as annuals and perennials?
(Q8) Biome mediates responses of arthropod communities to diversified farming.	Meta-regression: Do effect sizes differ among boreal, Mediterranean, temperate, and tropical biomes?

**Table S8.** Results of one-sample *t*-tests testing whether organic farming and in-field plant diversification impacted overall arthropod communities (pooled across functional groups). Means are average untransformed log-response ratios comparing organic to conventional, or high to low in-field plant diversity, data. Effect sizes transformed to percent change are in parentheses.

Management scheme	Metric	N	Mean	SE	<i>t</i>	<i>p</i> -value
Organic vs. conventional	Abundance	81	0.36 (45%)	0.10	3.76	0.0003
Organic vs. conventional	Local richness	81	0.19 (21%)	0.05	3.75	0.0003
Organic vs. conventional	Local evenness	73	-0.06 (-6%)	0.04	-1.69	0.095
Organic vs. conventional	Regional richness	81	0.11 (10%)	0.04	2.52	0.014
Organic vs. conventional	Regional evenness	81	-0.08 (-9%)	0.04	-1.87	0.065
In-field plant diversity	Abundance	29	0.24 (27%)	0.16	1.48	0.15
In-field plant diversity	Local richness	29	0.21 (23%)	0.08	2.49	0.019
In-field plant diversity	Local evenness	26	-0.07 (-6%)	0.05	-1.31	0.20
In-field plant diversity	Regional richness	29	0.17 (19%)	0.08	2.24	0.033
In-field plant diversity	Regional evenness	29	-0.14 (-13%)	0.09	-1.51	0.14

**Table S9.** Best-fit models, with  $\Delta AICc < 2$ , and global models explaining arthropod abundance, richness, and evenness in fields managed organically vs. conventionally.  $K$  is the number of estimated model parameters (fixed plus random effects). Parameters are: F=functional group, D=diversity scale (local, regional), LC=landscape complexity (simple, complex), A=cultivation period (annual, perennial), B=biome. A “\*” indicates an interaction and both of its main effects. Detritivores were excluded from meta-regressions due to low sample size.

Abundance					
Model ID	Parameters	$K$	AICc	$\Delta AICc$	weight
2	A	4	178.1	0	0.40
6	F + A	6	178.6	0.41	0.32
14	F + A + LC	7	178.8	0.69	0.28
Global	F×D + F×LC + D×LC + A + B	12	191.4	13.26	
Richness					
Model ID	Parameters	$K$	AICc	$\Delta AICc$	weight
61	D + F×LC	9	148.1	0	0.57
45	F×LC	8	148.6	0.54	0.43
Global	F×D + F×LC + D×LC + A + B	16	163.2	15.1	
Evenness					
Model ID	Parameters	$K$	AICc	$\Delta AICc$	weight
1	intercept only	3	82.5	0	0.52
17	D	4	84.0	1.5	0.25
2	A	4	84.0	1.5	0.24
Global	F×D + F×LC + D×LC + A + B	16	102.7	20.2	

**Table S10.** Regression details for best-fit models listed in Table S7 that explain arthropod abundance, richness, and evenness in fields managed organically vs. conventionally. We significance of fixed effects with likelihood ratio tests (LRTs), and used post-hoc planned contrasts (with *p*-values adjusted via Holm's sequential Bonferroni procedure) to test for (1) differences in effect size among functional groups, and (2) differences in effect size between the local and regional scales within each functional group. Parameters are: F=functional group (h=herbivore, po=pollinator, pr=predator), D=diversity scale (r=regional), LC= landscape complexity (c=complex, s=simple), A=cultivation period (p=perennial), B=biome (b=boreal, M=Mediterranean, te=temperate, tr=tropical). A “:” indicates an interaction. Detritivores were excluded from meta-regressions due to low sample size.

<b>Abundance (detritivores excluded)</b>									
<b>Model ID</b>	<b>Parameter</b>	<b>Coefficient (SE)</b>	<b>LRT <math>\chi^2</math></b>	<b>LRT df</b>	<b>LRT <i>p</i>-value</b>	<b>Contrast</b>	<b>Contrast <math>\chi^2</math></b>	<b>Contrast df</b>	<b>Contrast <i>p</i>-value</b>
2	Intercept	0.54 (0.13)	NA						
	A, p	-0.50 (0.24)	4.48	1	0.034				
6	Intercept	0.41 (0.24)	NA			F, h-po	2.96	1	0.18
	F, po	0.52 (0.30)	4.36	2	0.11	F, h-pr	0.01	1	0.91
	F, pr	0.03 (0.28)				F, po-pr	3.51	1	0.18
	A, p	-0.62 (0.24)	6.11	1	0.014				
14	Intercept	0.09 (0.33)	NA			F, h-po	4.87	1	0.075
	F, po	0.75 (0.34)	6.41	2	0.041	F, h-pr	0.23	1	0.63
	F, pr	0.14 (0.28)				F, po-pr	5.04	1	0.074
	LC, s	0.36 (0.25)	2.22	1	0.14				
	A, p	-0.57 (0.24)	5.68	1	0.017				
<b>Richness (detritivores excluded)</b>									
<b>Model ID</b>	<b>Parameter</b>	<b>Coefficient (SE)</b>	<b>LRT <math>\chi^2</math></b>	<b>LRT df</b>	<b>LRT <i>p</i>-value</b>	<b>Contrast</b>	<b>Contrast <math>\chi^2</math></b>	<b>Contrast df</b>	<b>Contrast <i>p</i>-value</b>
61	Intercept	-0.46 (0.21)	NA			F, h-po	10.23	1	0.004
	F, po	0.88 (0.20)	18.46	4	0.001	F, h-pr	8.14	1	0.009
	F, pr	0.68 (0.20)				F, po-pr	1.81	1	0.18
	S, r	-0.09 (0.06)	2.85	1	0.092	F:LC, c-s in h	6.88	1	0.026
	LC, s	0.61 (0.23)	10.66	3	0.014	F:LC, c-s in po	0.31	1	1
	F:LC, po	-0.75 (0.32)	10.64	2	0.005	F:LC, c-s	0.42	1	1



						in pr			
	F:LC, pr	-0.72 (0.22)							
45	Intercept	-0.51 (0.21)	NA			F, h-po	10.13	1	0.004
	F, po	0.88 (0.20)	17.95	4	0.001	F, h-pr	7.94	1	0.010
	F, pr	0.68 (0.21)				F, po-pr	1.82	1	0.18
	LC, s	0.61 (0.24)	10.30	3	0.016	F:LC, c-s in h	6.77	1	0.028
	F:LC, po:s	-0.75 (0.32)	10.27	2	0.006	F:LC, c-s in po	0.32	1	1
	F:LC, pr:s	-0.72 (0.23)				F:LC, c-s in pr	0.41	1	1
Evenness (detritivores excluded)									
Model ID	Parameter	Coefficient (SE)	LRT $\chi^2$	LRT df	LRT <i>p</i> - value				
17	Intercept	-0.08 (0.05)	NA						
	S, r	-0.04 (0.05)	0.65	1	0.42				
2	Intercept	-0.12 (0.06)	NA						
	A, p	0.07 (0.10)	0.61	1	0.43				

**Table S11.** Best-fit models, with  $\Delta\text{AICc} < 2$ , and global models explaining arthropod abundance, richness, and evenness in fields managed with high vs. low in-field plant diversity.  $K$  is the number of estimated model parameters (fixed plus random effects). Parameters are: F=functional group, D=diversity scale (local, regional), LC=landscape complexity (simple, complex), A=cultivation period (annual, perennial), B=biome. A “\*” indicates an interaction and both of its main effects.

<b>Abundance</b>					
<b>Model ID</b>	<b>Parameters</b>	<b><math>K</math></b>	<b>AICc</b>	<b><math>\Delta\text{AICc}</math></b>	<b>weight</b>
1	intercept only	3	70.4	0	0.67
2	A	4	71.7	1.4	0.33
Global	F×D + F×LC + D×LC + A + B	14	96.7	26.3	
<b>Richness</b>					
<b>Model ID</b>	<b>Parameters</b>	<b><math>K</math></b>	<b>AICc</b>	<b><math>\Delta\text{AICc}</math></b>	<b>weight</b>
5	F	6	42.2	0	0.36
45	F×LC	10	42.2	0.04	0.36
7	F + B	9	42.7	0.5	0.28
Global	F×D + F×LC + D×LC + A + B	19	54.1	11.9	
<b>Evenness</b>					
<b>Model ID</b>	<b>Parameters</b>	<b><math>K</math></b>	<b>AICc</b>	<b><math>\Delta\text{AICc}</math></b>	<b>weight</b>
85	F×D	10	21.8	0	1
Global	F×D + F×LC + D×LC + A + B	19	48.5	26.7	

107 **Table S12.** Regression details for best-fit models listed in Table S9 that explain arthropod abundance, richness, and evenness in fields  
108 managed with high vs. low in-field plant diversity. We significance of fixed effects with likelihood ratio tests (LRTs), and used post-  
109 hoc planned contrasts (with *p*-values adjusted via Holm’s sequential Bonferroni procedure) to test for (1) differences in effect size  
110 among functional groups, (2) differences in effect size between the local and regional scales within each functional group, and (3)  
111 landscape complexity differences among each pair of functional groups. Parameters are: F=functional group (d=detritivore,  
112 h=herbivore, po=pollinator, pr=predator), D=diversity scale (l=local, r=regional), LC= landscape complexity (c=complex, s=simple),  
113 A=cultivation period (p=perennial), B=biome (b=boreal, M=Mediterranean, te=temperate, tr=tropical). A “:” indicates an interaction.

Abundance									
Model ID	Parameter	Coefficient (SE)	LRT $\chi^2$	LRT df	LRT <i>p</i> -value				
2	Intercept	0.06 (0.20)	NA						
	A, p	0.40 (0.36)	1.33	1	0.25				
Richness									
Model ID	Parameter	Coefficient (SE)	LRT $\chi^2$	LRT df	LRT <i>p</i> -value	Contrast	Contrast $\chi^2$	Contrast df	Contrast <i>p</i> -value
5	Intercept	0.25 (0.16)	NA			F, d-h	6.24	1	0.075
	F, h	-0.30 (0.12)	9.57	3	0.023	F, d-po	0.10	1	1
	F, po	0.06 (0.19)				F, d-pr	4.13	1	0.21
	F, pr	-0.24 (0.12)				F, h-po	4.02	1	0.21
						F, h-pr	0.31	1	1
						F, po-pr	3.17	1	0.23
45	Intercept	0.19 (0.20)	NA			F, d-h	10.37	1	0.008
	F, h	-0.03 (0.14)	20.36	6	0.002	F, d-po	0.07	1	1
	F, po	0.19 (0.25)				F, d-pr	7.16	1	0.037
	F, pr	-0.21 (0.14)				F, h-po	2.74	1	0.39
	LC, s	0.32 (0.34)	11.00	4	0.027	F, h-pr	0.43	1	1
	F:LC, h:s	-0.67 (0.23)	10.57	3	0.014	F, po-pr	1.82	1	0.53
	F:LC, po:s	-0.49 (0.40)				F:LC, c-s in d	0.93	1	1
	F:LC, pr:s	-0.18 (0.23)				F:LC, c-s in h	1.28	1	1
						F:LC, c-s in po	0.52	1	1

						F:LC, c-s in pr	0.24	1	1
7	Intercept	0.30 (0.38)	NA			F, d-h	6.54	1	0.064
	F, h	-0.31 (0.12)	11.30	3	0.010	F, d-po	0.29	1	0.84
	F, po	0.10 (0.18)				F, d-pr	3.49	1	0.19
	F, pr	-0.23 (0.12)				F, h-po	5.67	1	0.086
	B, M	0.17 (0.40)	7.61	3	0.054	F, h-pr	0.65	1	0.84
	B, te	-0.28 (0.39)				F, po-pr	3.93	1	0.19
	B, tr	0.09 (0.41)				B, b-M	0.18	1	1
						B, b-te	0.51	1	1
						B, b-tr	0.05	1	1
						B, M-te	5.54	1	1
						B, M-tr	0.14	1	1
						B, te-tr	3.56	1	1
<b>Richness, boreal data excluded</b>									
Model ID	Parameter	Coefficient (SE)	LRT $\chi^2$	LRT df	LRT <i>p</i> - value	Contrast	Contrast $\chi^2$	Contrast df	Contrast <i>p</i> -value
7	Intercept	0.47 (0.20)	NA			F, d-h	6.36	1	0.070
	F, h	-0.31 (0.12)	10.90	3	0.012	F, d-po	0.29	1	0.85
	F, po	0.10 (0.19)				F, d-pr	3.40	1	0.20
	F, pr	-0.23 (0.12)				F, h-po	5.45	1	0.087
	B, te	-0.45 (0.19)	7.23	2	0.027	F, h-pr	0.64	1	0.85
	B, tr	-0.08 (0.22)				F, po-pr	3.92	1	0.19
						B, M-te	5.54	1	0.056
						B, M-tr	0.14	1	0.71
						B, te-tr	3.56	1	0.12
47	Intercept	0.41 (0.22)	NA			F, d-h	10.56	1	0.007
	F, h	-0.03 (0.14)	21.29	6	0.002	F, d-po	0.01	1	0.95
	F, po	0.18 (0.27)				F, d-pr	6.55	1	0.052
	F, pr	-0.19 (0.14)				F, h-po	4.04	1	0.18
	LC, s	0.31 (0.36)	10.55	4	0.032	F, h-pr	0.68	1	0.82
	B, te	-0.43 (0.21)	7.07	2	0.029	F, po-pr	2.55	1	0.33

	B, tr	-0.05 (0.29)				B, M-te	4.39	1	0.11
	F:LC, h:s	-0.69 (0.23)	10.30	3	0.016	B, M-tr	0.03	1	0.86
	F:LC, po:s	-0.39 (0.47)				B, te-tr	2.40	1	0.24
	F:LC, pr:s	-0.22 (0.23)				F:LC, c-s in d	0.73	1	1
						F:LC, c-s in h	1.23	1	1
						F:LC, c-s in po	0.12	1	1
						F:LC, c-s in pr	0.08	1	1
5	Intercept	0.24 (0.17)	NA			F, d-h	6.04	1	0.084
	F, h	-0.30 (0.12)	9.21	3	0.027	F, d-po	0.12	1	1
	F, po	0.07 (0.20)				F, d-pr	4.02	1	0.22
	F, pr	-0.25 (0.12)				F, h-po	3.84	1	0.22
						F, h-pr	0.29	1	1
						F, po-pr	2.98	1	0.25
Evenness									
Model ID	Parameter	Coefficient (SE)	LRT $\chi^2$	LRT df	LRT <i>p</i> -value	Contrast	Contrast $\chi^2$	Contrast df	Contrast <i>p</i> -value
85	Intercept	-0.08 (0.21)	NA			F, d-h	17.99	1	0.0001
	F, h	0.14 (0.21)	46.79	6	<0.0001	F, d-po	6.45	1	0.045
	F, po	-0.04 (0.23)				F, d-pr	21.60	1	<0.0001
	F, pr	0.13 (0.20)				F, h-po	0.59	1	0.89
	S, r	-0.88 (0.21)	16.44	4	0.003	F, h-pr	0.18	1	0.89
	F:S, h:r	0.79 (0.24)	16.13	3	0.001	F, po-pr	1.21	1	0.81
	F:S, po:r	0.92 (0.22)				F:S, l-r in d	17.44	1	0.0001
	F:S, pr:r	0.89 (0.23)				F:S, l-r in h	0.55	1	1
						F:S, l-r in po	0.44	1	1
						F:S, l-r in pr	0.01	1	1

**Table S13.** Results of one-sample *t*-tests testing whether organic farming and in-field plant diversification impacted overall arthropod communities (pooled across functional groups) for rare and common taxa. We classified taxa as common if their relative abundance was at least 5% of the total community; other species were categorized as rare. Means are average untransformed log-response ratios comparing organic to conventional, or high to low in-field plant diversity, data. Effect sizes transformed to percent change are in parentheses.

Management scheme	Metric	Relative abundance category	N	Mean	SE	<i>t</i>	<i>p</i> -value
Organic vs. conventional	Abundance	Rare	77	0.44 (55%)	0.45	4.16	<0.0001
Organic vs. conventional	Abundance	Common	82	0.37 (45%)	0.51	3.64	<0.0001
Organic vs. conventional	Local richness	Rare	77	0.24 (27%)	0.38	3.29	0.002
Organic vs. conventional	Local richness	Common	82	0.13 (14%)	0.31	2.75	0.007
Organic vs. conventional	Regional richness	Rare	73	0.12 (12%)	0.31	2.52	0.014
Organic vs. conventional	Regional richness	Common	78	0.05 (6%)	0.29	1.80	0.076
In-field plant diversity	Abundance	Rare	25	0.23 (25%)	1.31	1.33	0.19
In-field plant diversity	Abundance	Common	30	0.31 (37%)	1.10	1.79	0.084
In-field plant diversity	Local richness	Rare	25	0.33 (39%)	0.68	2.24	0.035
In-field plant diversity	Local richness	Common	30	0.13 (14%)	0.31	2.17	0.038
In-field plant diversity	Regional richness	Rare	24	0.24 (28%)	0.69	1.89	0.071
In-field plant diversity	Regional richness	Common	25	0.04 (4%)	0.18	1.45	0.16

122 **Table S14.** Results of paired *t*-tests testing whether organic farming and in-field plant diversification impacted arthropod abundance  
123 and richness differentially for rare and common taxa. Means are average untransformed log-response ratios comparing organic to  
124 conventional, or high to low in-field plant diversity, data. Effect sizes transformed to percent change are in parentheses.  
125

Management scheme	Metric	N common taxa	Mean common taxa	SE common taxa	N rare taxa	Mean rare taxa	SE rare taxa	<i>t</i>	<i>p</i> -value
Organic vs. conventional	Abundance	82	0.37 (45%)	0.10	77	0.44 (55%)	0.11	-0.76	0.45
Organic vs. conventional	Local richness	82	0.13 (14%)	0.05	77	0.24 (27%)	0.07	-2.40	0.019
Organic vs. conventional	Regional richness	78	0.05 (6%)	0.03	73	0.12 (12%)	0.05	-1.63	0.11
In-field plant diversity	Abundance	30	0.31 (37%)	0.17	25	0.23 (25%)	0.17	1.02	0.32
In-field plant diversity	Local richness	30	0.13 (14%)	0.06	25	0.33 (39%)	0.15	-1.61	0.12
In-field plant diversity	Regional richness	25	0.04 (4%)	0.02	24	0.24 (28%)	0.13	-1.48	0.15

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**Table S15.** Results of one-sample *t*-tests testing whether organic farming and in-field plant diversification impacted pollinator communities. Means are average untransformed log-response ratios comparing organic to conventional, or high to low in-field plant diversity, data. Effect sizes transformed to percent change are in parentheses.

Management scheme	Metric	N	Mean	SE	<i>t</i>	<i>p</i> -value
Organic vs. conventional	Abundance	20	0.61 (90%)	0.30	2.01	0.058
Organic vs. conventional	Local richness	20	0.42 (55%)	0.16	2.68	0.015
Organic vs. conventional	Local evenness	17	0.05 (5%)	0.10	0.52	0.61
Organic vs. conventional	Regional richness	20	0.27 (32%)	0.12	2.25	0.036
Organic vs. conventional	Regional evenness	20	-0.15 (-15%)	0.10	-1.58	0.13
In-field plant diversity	Abundance	13	0.37 (45%)	0.14	2.62	0.023
In-field plant diversity	Local richness	13	0.36 (44%)	0.12	2.97	0.012
In-field plant diversity	Local evenness	13	-0.11 (-11%)	0.05	-2.07	0.061
In-field plant diversity	Regional richness	13	0.25 (29%)	0.13	2.01	0.068
In-field plant diversity	Regional evenness	13	-0.07 (-6%)	0.13	-0.51	0.62



**Table S16.** Results of one-sample *t*-tests testing whether organic farming and in-field plant diversification impacted predator communities. Means are average untransformed log-response ratios comparing organic to conventional, or high to low in-field plant diversity, data. Effect sizes transformed to percent change are in parentheses.

Management scheme	Metric	N	Mean	SE	<i>t</i>	<i>p</i> -value
Organic vs. conventional	Abundance	36	0.32 (39%)	0.09	3.41	0.0020
Organic vs. conventional	Local richness	36	0.15 (14%)	0.06	2.42	0.021
Organic vs. conventional	Local evenness	35	-0.09 (-9%)	0.03	-2.69	0.011
Organic vs. conventional	Regional richness	36	0.09 (6%)	0.06	1.50	0.14
Organic vs. conventional	Regional evenness	36	-0.12 (-14%)	0.05	-2.35	0.024
In-field plant diversity	Abundance	8	-0.10 (-10%)	0.34	-0.29	0.78
In-field plant diversity	Local richness	8	-0.03 (-3%)	0.13	-0.19	0.85
In-field plant diversity	Local evenness	8	0.06 (6%)	0.10	0.54	0.61
In-field plant diversity	Regional richness	8	0.05 (5%)	0.10	0.51	0.63
In-field plant diversity	Regional evenness	8	0.07 (7%)	0.10	0.63	0.55

**Table S17.** Results of one-sample *t*-tests testing whether organic farming and in-field plant diversification impacted herbivore communities. Means are average untransformed log-response ratios comparing organic to conventional, or high to low in-field plant diversity, data. Effect sizes transformed to percent change are in parentheses.

Management scheme	Metric	N	Mean	SE	<i>t</i>	<i>p</i> -value
Organic vs. conventional	Abundance	17	0.24 (23%)	0.18	1.30	0.21
Organic vs. conventional	Local richness	17	0.12 (10%)	0.08	1.44	0.17
Organic vs. conventional	Local evenness	14	-0.16 (-14%)	0.12	-1.33	0.21
Organic vs. conventional	Regional richness	17	0.07 (5%)	0.07	1.06	0.30
Organic vs. conventional	Regional evenness	17	-0.07 (-7%)	0.08	-0.89	0.39
In-field plant diversity	Abundance	5	0.25 (28%)	0.42	0.58	0.59
In-field plant diversity	Local richness	5	0.17 (18%)	0.25	0.67	0.54
In-field plant diversity	Local evenness	4	-0.04 (-4%)	0.12	-0.30	0.78
In-field plant diversity	Regional richness	5	-0.04 (-4%)	0.23	-0.20	0.85
In-field plant diversity	Regional evenness	5	-0.10 (-10%)	0.15	0.68	0.53

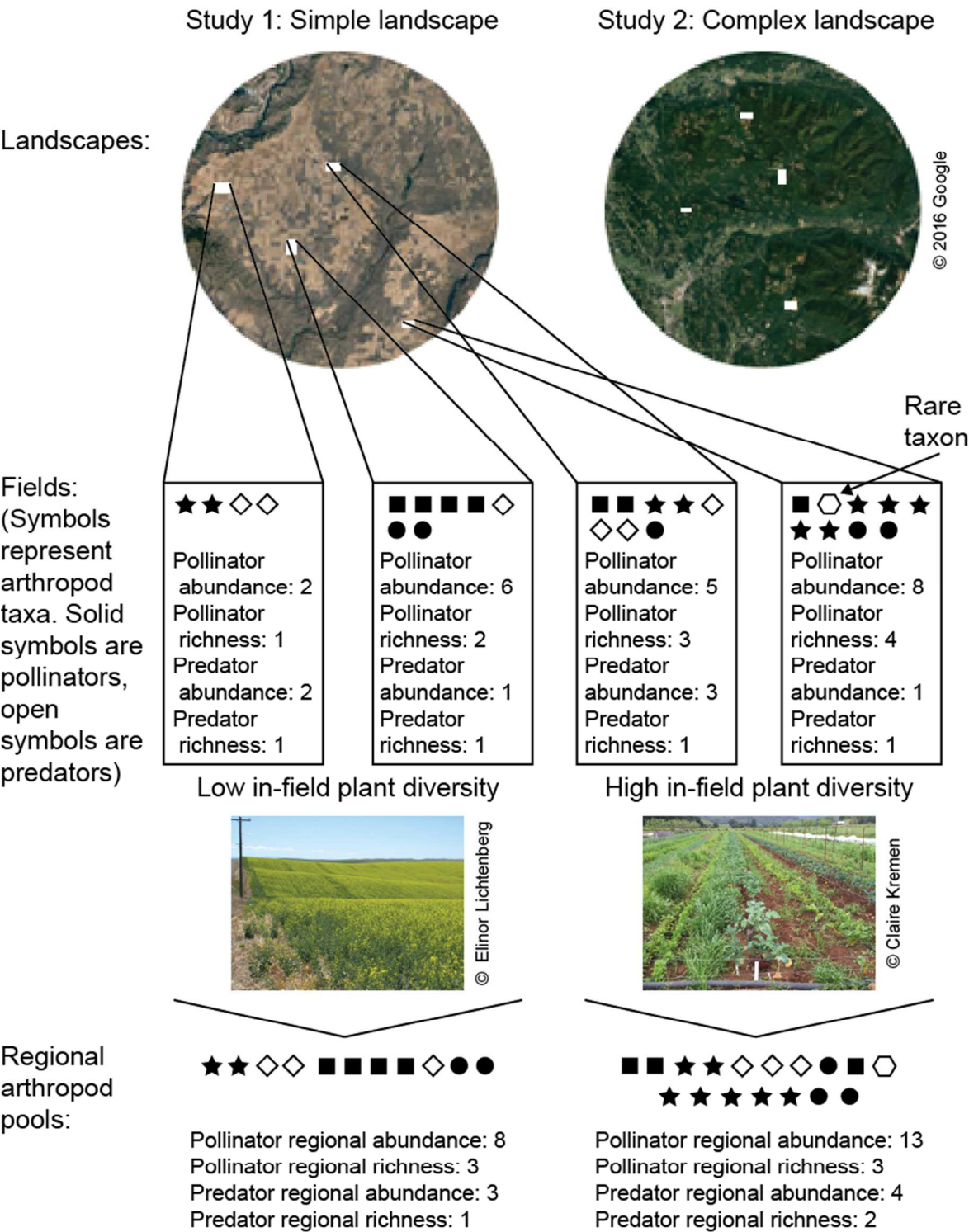
**Table S18.** Results of one-sample *t*-tests testing whether organic farming and in-field plant diversification impacted detritivore communities. Means are average untransformed log-response ratios comparing organic to conventional, or high to low in-field plant diversity, data. Effect sizes transformed to percent change are in parentheses.

Management scheme	Metric	N	Mean	SE	<i>t</i>	<i>p</i> -value
Organic vs. conventional	Abundance	8	0.23 (26%)	0.29	0.79	0.46
Organic vs. conventional	Local richness	8	0.01 (1%)	0.07	0.15	0.89
Organic vs. conventional	Local evenness	7	-0.01 (-1%)	0.09	-0.06	0.95
Organic vs. conventional	Regional richness	8	-0.10 (-9%)	0.11	-0.91	0.39
Organic vs. conventional	Regional evenness	8	0.26 (30%)	0.21	1.28	0.24
In-field plant diversity	Abundance	3	0.55 (74%)	1.03	0.54	0.65
In-field plant diversity	Local richness	3	0.25 (28%)	0.31	0.82	0.50
In-field plant diversity	Local evenness	1	-0.57 (-44%)	NA	NA	NA
In-field plant diversity	Regional richness	3	0.52 (69%)	0.07	7.51	0.017
In-field plant diversity	Regional evenness	3	-1.06 (-65%)	0.22	-4.80	0.041

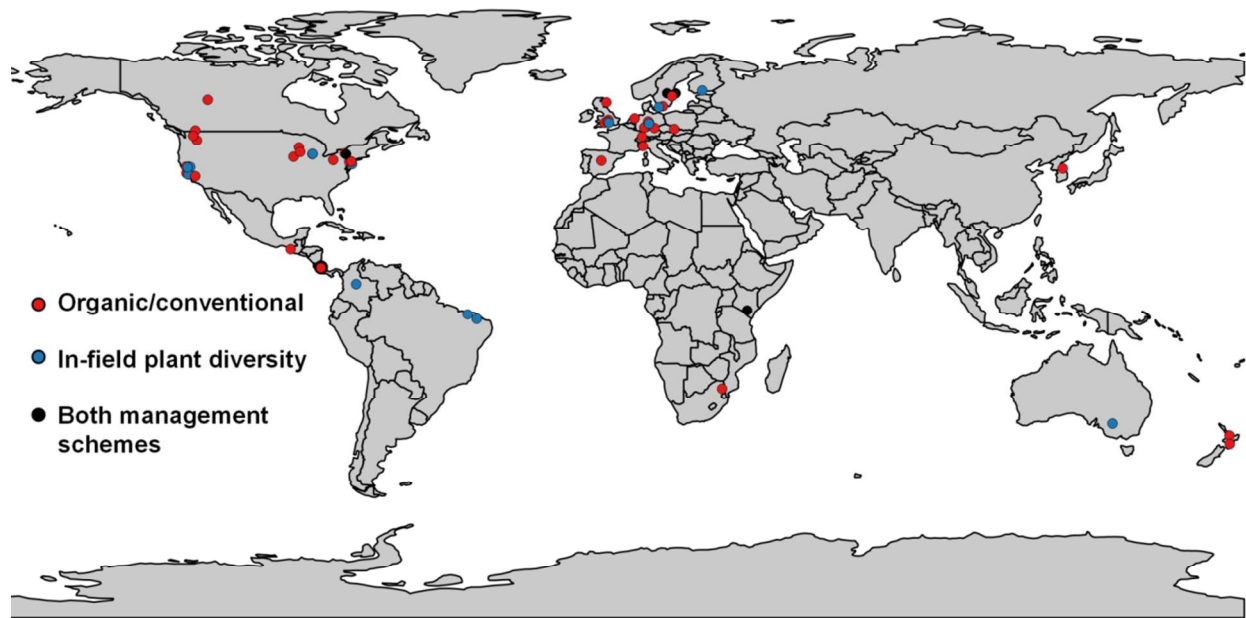
**Table S19.** Results of paired *t*-tests testing whether organic farming and in-field plant diversification impacted arthropod abundance and richness differentially for rare and common taxa, in simple and complex landscapes. Means are average untransformed log-response ratios comparing organic to conventional, or high to low in-field plant diversity, data. Effect sizes transformed to percent change are in parentheses.

Management scheme	Metric	Landscape complexity	N common taxa	Mean common taxa	SE common taxa	N rare taxa	Mean rare taxa	SE rare taxa	<i>t</i>	<i>p</i> -value
Organic vs. conventional	Abundance	Simple	45	0.45 (57%)	0.12	43	0.36 (44%)	0.11	0.51	0.61
Organic vs. conventional	Abundance	Complex	30	0.28 (33%)	0.21	28	0.58 (78%)	0.24	-1.90	0.068
Organic vs. conventional	Local richness	Simple	45	0.09 (10%)	0.05	43	0.15 (16%)	0.07	-0.88	0.39
Organic vs. conventional	Local richness	Complex	30	0.19 (21%)	0.10	28	0.36 (44%)	0.16	-2.35	0.027
Organic vs. conventional	Regional richness	Simple	42	0.05 (6%)	0.04	41	0.06 (6%)	0.06	0.10	0.92
Organic vs. conventional	Regional richness	Complex	29	0.04 (4%)	0.04	26	0.16 (17%)	0.07	-2.33	0.028
In-field plant diversity	Abundance	Simple	13	0.24 (27%)	0.22	10	0.08 (8%)	0.07	1.58	0.15
In-field plant diversity	Abundance	Complex	17	0.37 (45%)	0.27	15	0.33 (39%)	0.28	0.05	0.96
In-field plant diversity	Local richness	Simple	13	0.09 (10%)	0.08	10	0.05 (5%)	0.10	1.00	0.35
In-field plant diversity	Local richness	Complex	17	0.16 (18%)	0.09	15	0.52 (68%)	0.23	-2.22	0.044
In-field plant diversity	Regional richness	Simple	10	0.06 (6%)	0.06	10	0.02 (2%)	0.09	-0.04	0.97
In-field plant diversity	Regional richness	Complex	15	0.02 (2%)	0.01	14	0.40 (50%)	0.20	-1.59	0.14

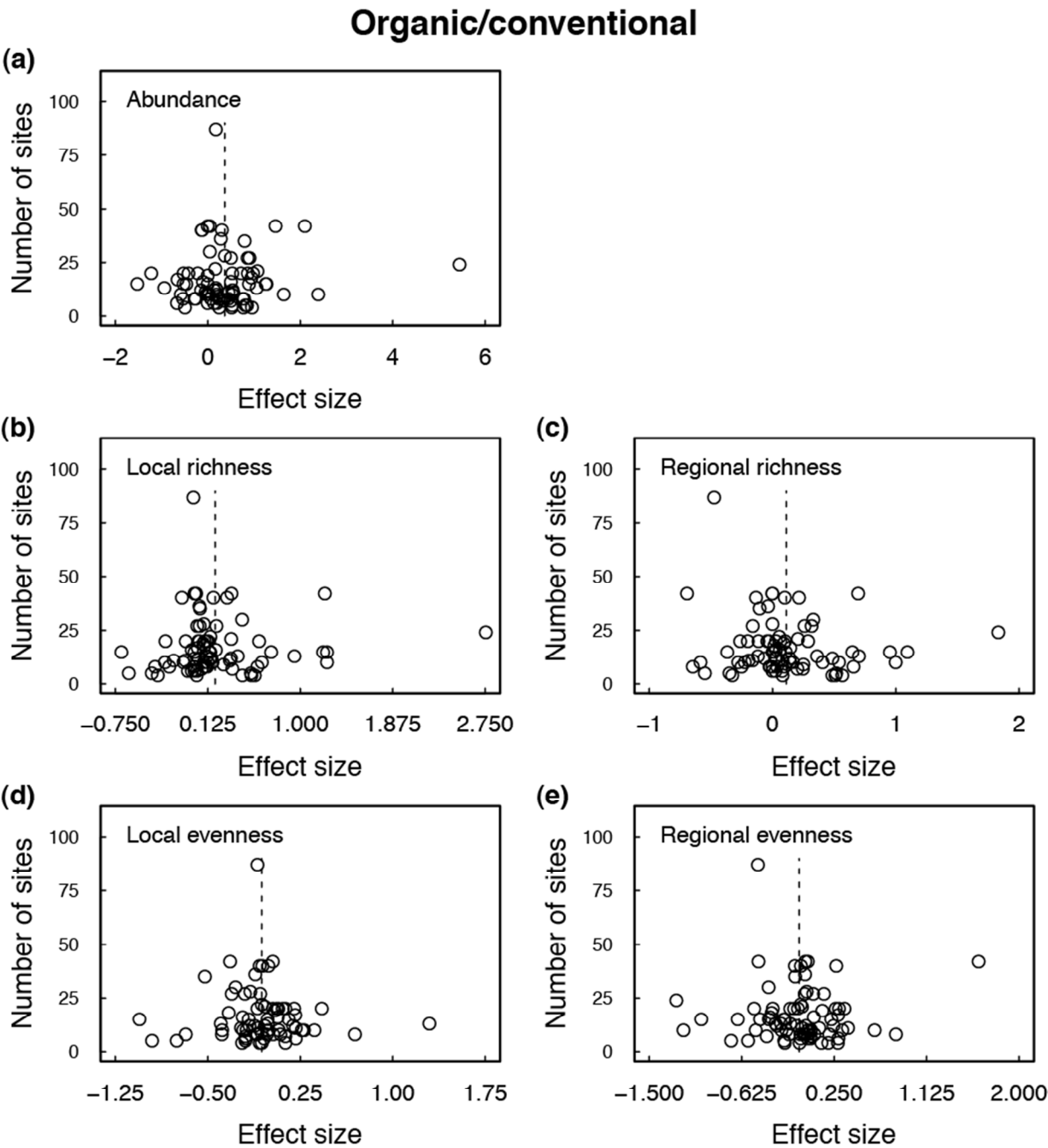
**Fig. S1.** Data structure and major factors used in the meta-analysis. Each study consisted of a collection of fields (white rectangles, not to scale) situated in simple or complex landscapes. We classified each field as having low or high in-field plant diversity, or being managed organically or conventionally (not shown). Within each study, we divided sampled taxa by functional group (detritivore, herbivore, pollinator, predator). For each sub-group, we calculated local abundance and diversity from field-level taxon pools, and regional diversity from the regional pool.



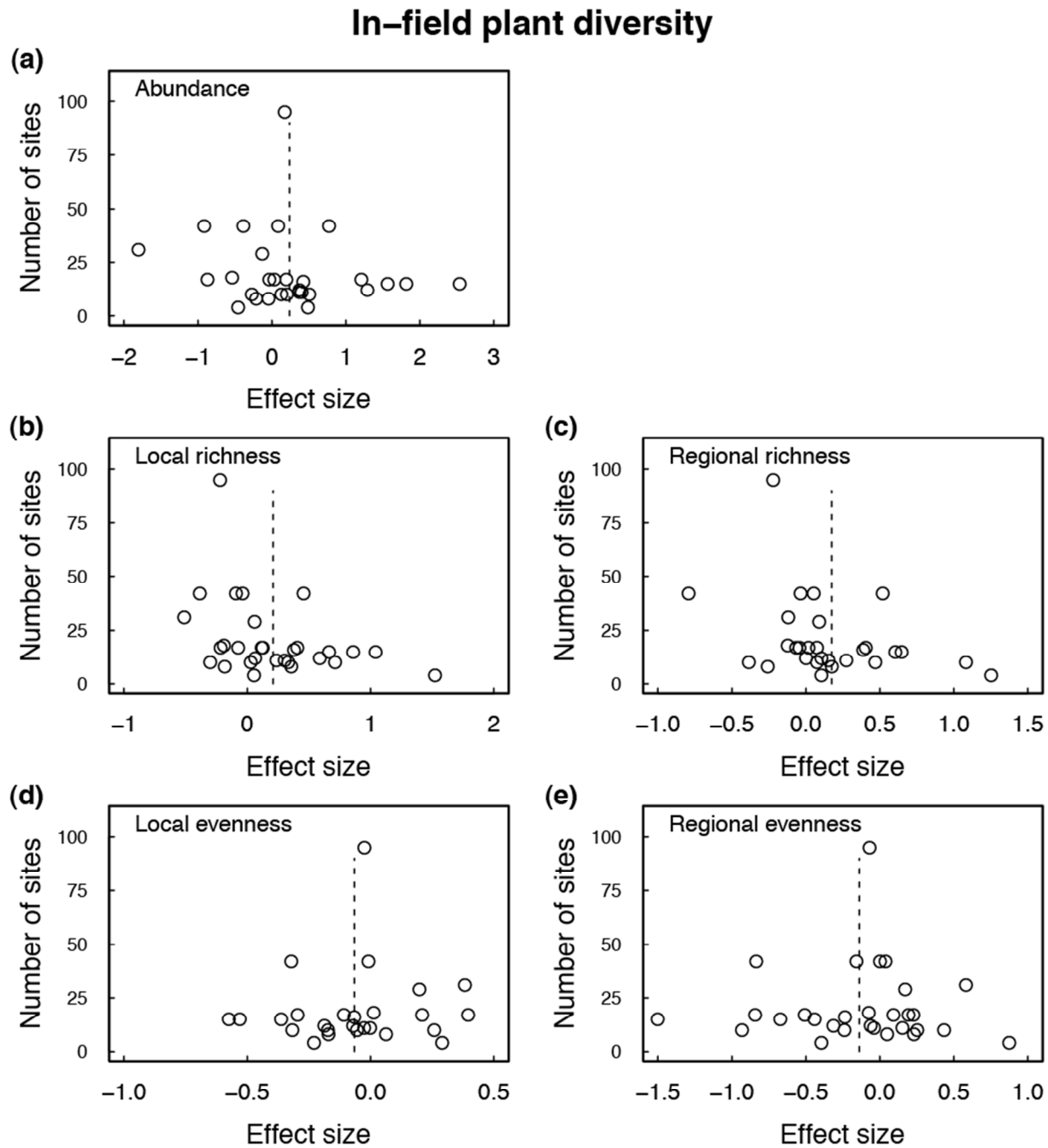
**Fig. S2.** Map of study sites.



**Figure S3.** Funnel plots for studies assessing organic vs. conventional farming. All plots are sufficiently symmetrical about their mean (visually assessed) to indicate no publication bias. Effect sizes are log-response ratios.

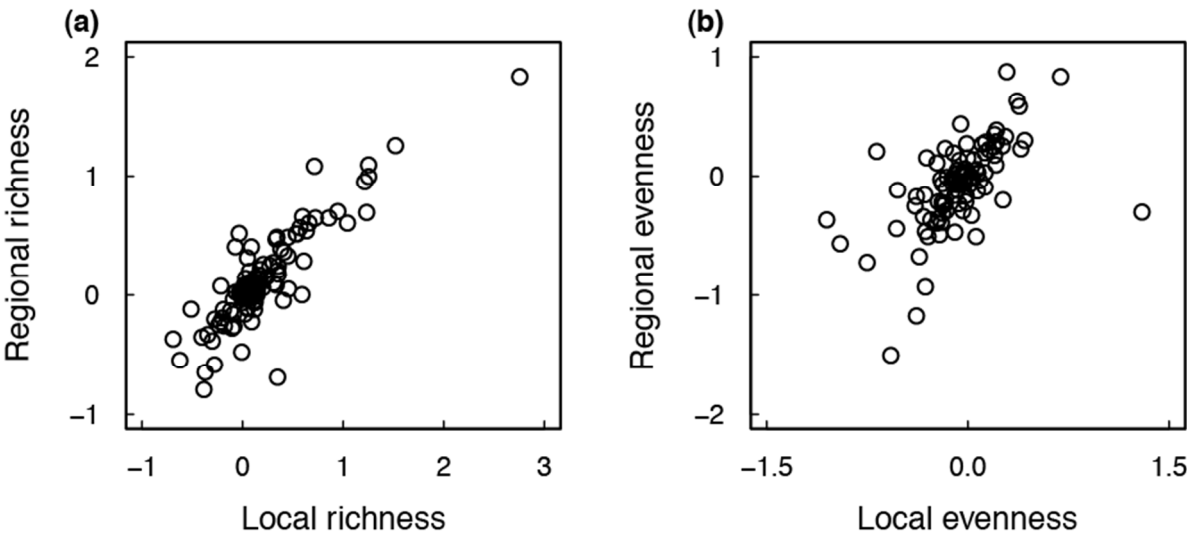


**Figure S4.** Funnel plots for studies assessing in-field plant diversification. All plots are sufficiently symmetrical about their mean (visually assessed) to indicate no publication bias. Effect sizes are log-response ratios.

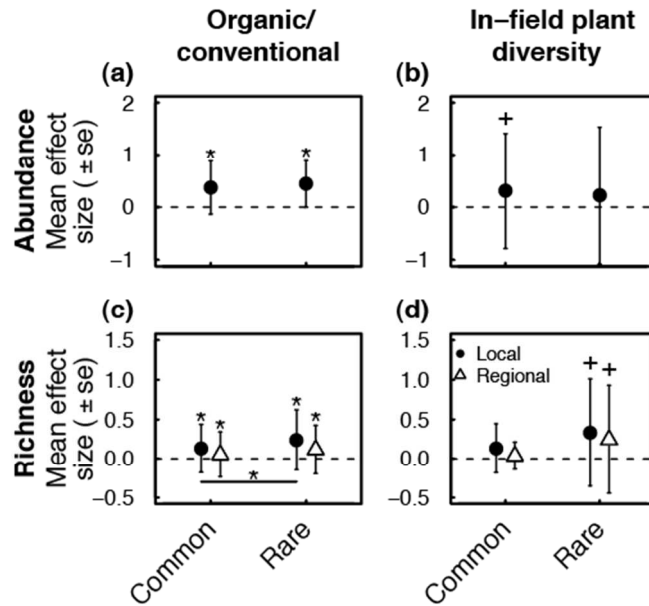




**Figure S5.** Diversity effects (log-response ratios) strongly correlated at the local and regional scales for both richness (Pearson’s correlation:  $r = 0.87$ ,  $t_{108} = 18.41$ ,  $p < 0.0001$ ) and evenness ( $r = 0.81$ ,  $t_{97} = 5.83$ ,  $p < 0.0001$ ).



**Figure S6.** Effects of farm management schemes on abundance (a, b) and richness (c, d) of common vs. rare taxa. Mean log-response ratios ( $\pm$ SE) of (left column) adopting organic farming and (right column) promoting in-field plant diversity. A “\*” ( $p < 0.05$ ) or “+” ( $0.05 \leq p < 0.1$ ) above a mean denotes a significant difference from zero (determined via one-sample  $t$ -tests), while one below a pair of means indicates a significant difference between rare and common taxa (determined via paired  $t$ -tests).



**Table S1.** Data holders and studies participating. We were unable to categorize landscape complexity when we obtained data directly from published articles that lacked GPS coordinates of sampling locations or information on natural habitat surrounding fields (Study IDs drit01, febe01, hesl01, hokk01, and weib01). These studies were excluded from meta-regressions.

Study ID	Reference or data holder	Crop(s)	Study location	Functional group(s)	Management scheme(s)	# sites (o=organic/ conventional, i-f=in-field plant diversity)	Year(s)
arms01	(Armstrong, 1995)	potato	Scotland	predators	organic/ conventional	4	1992
bata01	(Batáry <i>et al.</i> , 2012)	wheat	Germany	predators	organic/ conventional	18	2008
benj01	(Cariveau <i>et al.</i> , 2013)	blueberry	USA	pollinators	in-field plant diversity	16	2012
bere01	(Winqvist <i>et al.</i> , 2011)	wheat	Netherlands	predators	organic/ conventional	35	2007
bomm01	(Winqvist <i>et al.</i> , 2011)	wheat	Sweden	predators	organic/ conventional, in-field plant diversity	95	2007
bosq01	Bosque-Perez, Nilsa; Ramos, Mariangie	coffee	Costa Rica	herbivores	organic/ conventional, in-field plant diversity	18 (o), 19 (i-f)	2005
carv01	(Carvalho <i>et al.</i> , 2010, 2012)	mango	South Africa	herbivores, pollinators, predators	organic/ conventional	15	2009

chap01	(Chaplin-Kramer <i>et al.</i> , 2013)	broccoli	USA	detritivores, herbivores, predators	in-field plant diversity	17	2008
clou01	(Clough <i>et al.</i> , 2005, 2007a, 2007b)	wheat	Germany	detritivores, herbivores, predators	organic/ conventional, in-field plant diversity	42 (o), 17 (i-f)	2003
conn01	(Connelly <i>et al.</i> , 2015)	strawberry	USA	pollinators	organic/ conventional	13	2012
danf01	(Russo <i>et al.</i> , 2015)	apple	USA	pollinators	organic/ conventional, in-field plant diversity	10	2009
diek01	(Diekötter <i>et al.</i> , 2010)	wheat	Germany	detritivores, herbivores, predators	organic/ conventional	12	2007
drit01	(Dritschilo & Erwin, 1982)	corn	USA	predators	organic/ conventional	8	late 1970s?
eige01	Eigenbrode, Sanford	coffee	Costa Rica	predators	organic/ conventional	6	2001
ekro01	(Ekroos <i>et al.</i> , 2010)	various grains (combined)	Finland	predators	organic/ conventional, in-field plant diversity	28 (o), 29 (i-f)	1998
febe01	(Feber <i>et al.</i> , 1998)	wheat	England	predators	organic/ conventional	6	1995
frei01	Freitas, Breno	acerola	Brazil	pollinators	in-field plant diversity	4	2010

frei02	Freitas, Breno	cotton	Brazil	pollinators	in-field plant diversity	4	2010
fuku01	(Fukuda <i>et al.</i> , 2011)	pasture	New Zealand	detritivores, herbivores, predators	organic/ conventional	20	2009
gain01	Gaines, Hannah; Gratton, Claudio	cranberry	USA	pollinators	organic/ conventional	15	2008
hesl01	(Hesler <i>et al.</i> , 1993)	rice	USA	herbivores, predators	organic/ conventional	6	1988
hokk01	(Hokkanen & Holopainen, 1986)	cabbage	Germany	herbivores, predators	organic/ conventional	4	1982
holz01	(Holzschuh <i>et al.</i> , 2007)	wheat	Germany	pollinators	organic/ conventional, in-field plant diversity	42	2003
isaa01	(Isaacs & Kirk, 2010)	blueberry	USA	pollinators	in-field plant diversity	12	2008
isai01	(Isaia <i>et al.</i> , 2006)	grape	Italy	predators	organic/ conventional	5	2003
jha01	(Jha & Vandermeer, 2010)	coffee	Mexico	pollinators	organic/ conventional	7	2006
jona01	(Jonason <i>et al.</i> , 2013)	various grains (combined)	Sweden	herbivores, predators	organic/ conventional	36	2011

jone01	(Jones <i>et al.</i> , In press, In pressb; Mills <i>et al.</i> , In press)	apple	USA	herbivores, pollinators, predators	organic/ conventional	8	2011
klat01	Klatt, Björn; Tschardtke, Teja	strawberry	Germany	pollinators	in-field plant diversity	8	2010
klei01	Brittain, Claire; Klein, Alexandra	almond	USA	pollinators	organic/ conventional	13	2009
krau01	(Krauss <i>et al.</i> , 2011)	triticale	Germany	pollinators	organic/ conventional	24	2008
krem01	(Kremen <i>et al.</i> , 2002, 2004)	watermelon	USA	pollinators	organic/ conventional	21	2000
leto01	(Drinkwater <i>et al.</i> , 1995; Letourneau & Goldstein, 2001; Letourneau & Bothwell, 2007; Letourneau <i>et al.</i> , 2012, 2015)	broccoli, brussel sprouts	USA	predators	organic/ conventional, in-field plant diversity	10	2006
mall01	(Mallinger <i>et al.</i> , 2015)	apple	USA	pollinators	organic/ conventional	17	2012
mart01	(Martin <i>et al.</i> , 2016)	potato, daikon radish, rice, soybean	South Korea	predators	organic/ conventional	7 (radish), 8 (other crops)	2009

memm01	(Gibson <i>et al.</i> , 2007; Macfadyen <i>et al.</i> , 2009a, 2009b, 2011a, 2011b)	grains, brassicas, legumes	England	herbivores, predators	organic/ conventional	20 (grains), 5 (brassicas), 10 (legumes)	2005 (grains, legumes), 2006 (brassicas)
mora01	(Morandin & Winston, 2005, 2006)	canola	Canada	pollinators	organic/ conventional	16	2002
neam01	Elle, Elizabeth; Neame, Lisa	winter squash	Canada	pollinators	organic/ conventional	9	2010
ober01	(Öberg, 2007; Öberg <i>et al.</i> , 2007)	various grains (combined)	Sweden	predators	organic/ conventional, in-field plant diversity	8	2003 (i-f), 2004 (o)
otie01	(Otieno <i>et al.</i> , 2015)	pigeonpea	Kenya	pollinators	organic/ conventional, in-field plant diversity	12	2009
pfif01	(Pfiffner & Luka, 2003)	various grains (combined)	Switzerland	predators	organic/ conventional	12	1996-8
poco01	(Pocock & Jennings, 2008)	various grains (combined)	England	detritivores, herbivores, predators	organic/ conventional	40	2003
ponc01	(Ponce <i>et al.</i> , 2011)	wheat, barley	Spain	detritivores, herbivores, predators	organic/ conventional	27 (wheat), 11 (barley)	2008
pott01	(Carré <i>et al.</i> , 2009)	field bean	England	pollinators	in-field plant diversity	10	2005

pove01	(Poveda <i>et al.</i> , 2012); Martinez, Eliana	potato	Colombia	herbivores, predators	in-field plant diversity	11	2007
rose01	(de Valpine & Rosenheim, 2008)	cotton	USA	herbivores, predators	organic/ conventional	15	1993
rund01	(Bommarco <i>et al.</i> , 2012)	red clover	Sweden	pollinators	in-field plant diversity	17	2010
sard01	(Sardiñas & Kremen, 2015)	sunflower	USA	pollinators	in-field plant diversity	10	2011
saun01	(Saunders & Luck, 2013)	almond	Australia	detritivores, herbivores, predators	in-field plant diversity	15	2010
scho01	(Schon <i>et al.</i> , 2011)	pasture	New Zealand	detritivores, herbivores, predators	organic/ conventional	10	2007
scil01	Sciligo, Amber	strawberry	USA	pollinators	in-field plant diversity	17	2012
sidh01	(Sidhu, 2013)	squash	USA	pollinators	organic/ conventional	8	2011
snyd01	Crowder, David; Snyder, William	potato	USA	detritivores, herbivores, predators	organic/ conventional	20	2010
vese01	(Veselý & Šarapatka, 2008)	wheat, barley	Czech Republic	predators	organic/ conventional	4 (wheat), 4 (barley)	2001 (wheat), 2005 (barley)



weib01	(Weibull <i>et al.</i> , 2000)	cereals, clovers, grasses (combined)	Sweden	pollinators	organic/ conventional	16	1997-8
weis01	(Winqvist <i>et al.</i> , 2011)	wheat	Germany	predators	organic/ conventional, in-field plant diversity	30 (o), 31 (i-f)	2007
will01	Williams, Neal	watermelon	USA	pollinators	in-field plant diversity	10	2010
wils01	(Tuell <i>et al.</i> , 2009)	blueberry	USA	pollinators	organic/ conventional	15	2005
winf01	(Winfree <i>et al.</i> , 2007, 2008; Lonsdorf <i>et al.</i> , 2009; Rader <i>et al.</i> , 2013)	watermelon	USA	pollinators	organic/ conventional	10	2010
winf02	(Winfree <i>et al.</i> , 2008)	pepper, tomato	USA	pollinators	organic/ conventional	22 (pepper), 13 (tomato)	2004 (pepper), 2005 (tomato)

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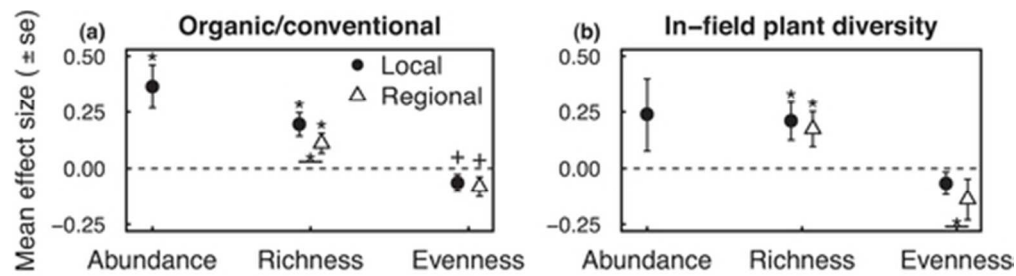
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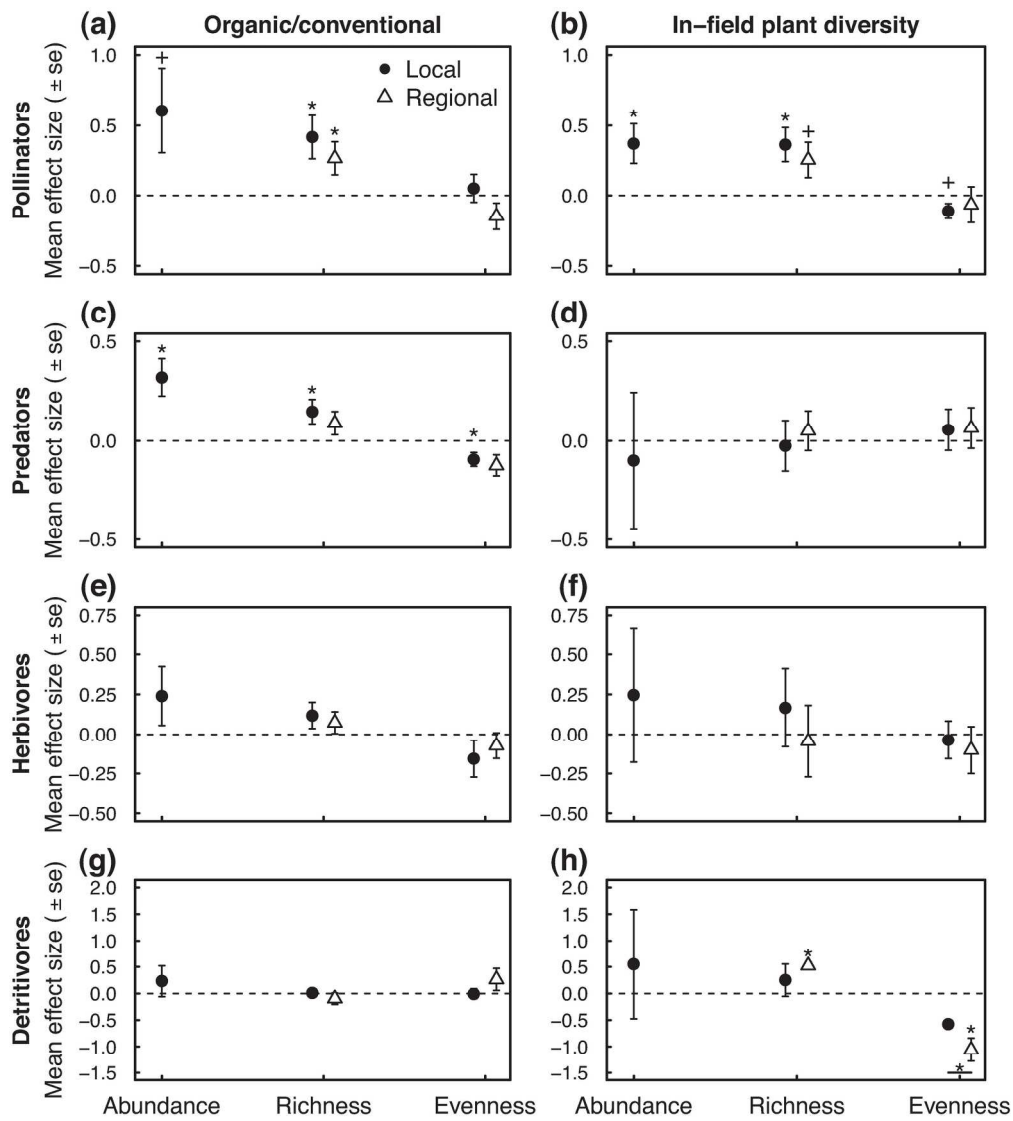
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Effects of farm management schemes on arthropod abundance, local diversity, and regional diversity. Values shown are for the entire arthropod community, and represent the mean log-response ratio ( $\pm$  SE) of (a) adopting organic farming and (b) promoting in-field plant diversity on abundance, richness, and evenness. A "\*" ( $p < 0.05$ ) or "+" ( $0.05 \leq p < 0.1$ ) above a mean denotes a significant difference from zero (determined via one-sample t-tests; statistical details in Table S8), while one below a pair of means indicates a significant difference between local and regional diversity (determined via linear mixed models; Tables S9-S12).

Fig. 1

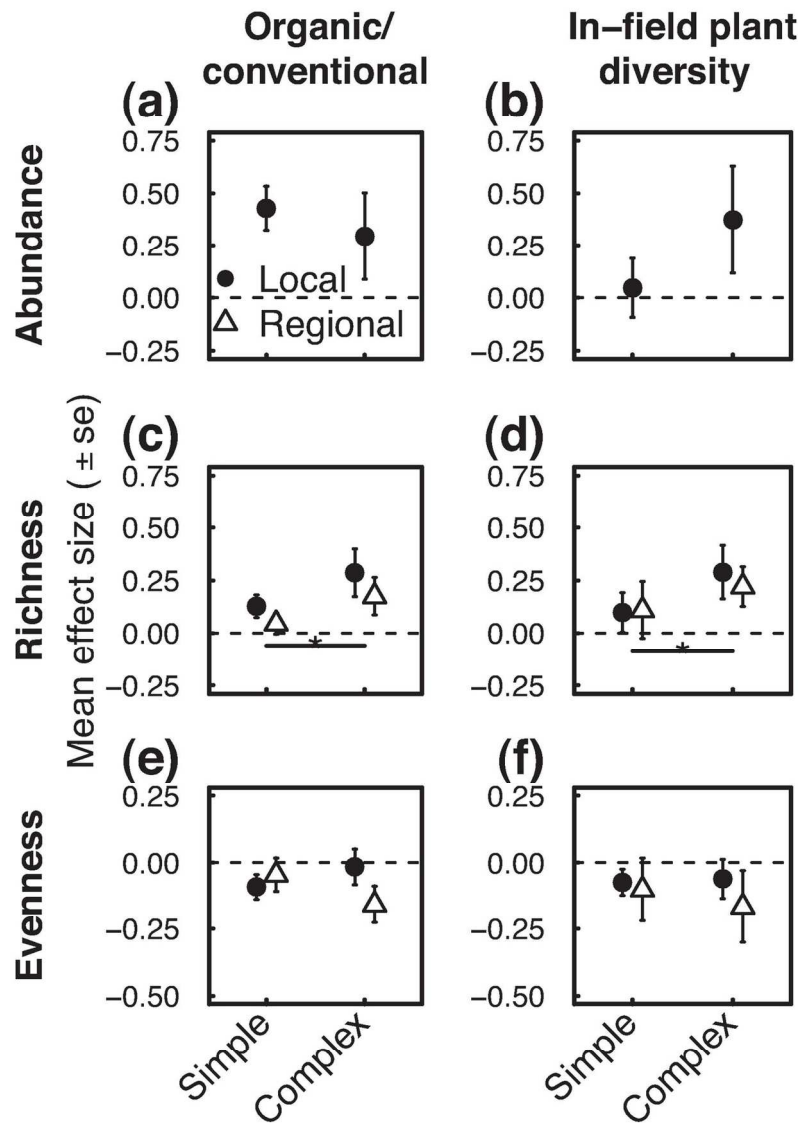
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Effects of farm management schemes on abundance, local diversity, and regional diversity of arthropod functional groups. Mean log-response ratios ( $\pm$  SE) of (left column) adopting organic farming and (right column) promoting in-field plant diversity for (a-b) pollinators, (c-d) predators, (e-f) herbivores, and (g-h) detritivores. A "\*" ( $p < 0.05$ ) or "+" ( $0.05 \leq p < 0.1$ ) above a mean denotes a significant difference from zero (determined via one-sample t-tests; Tables S15-S18). Meta-regressions indicated that differences between local and regional values did not vary with functional group (Tables S9-S12).

Fig. 2  
190x218mm (300 x 300 DPI)



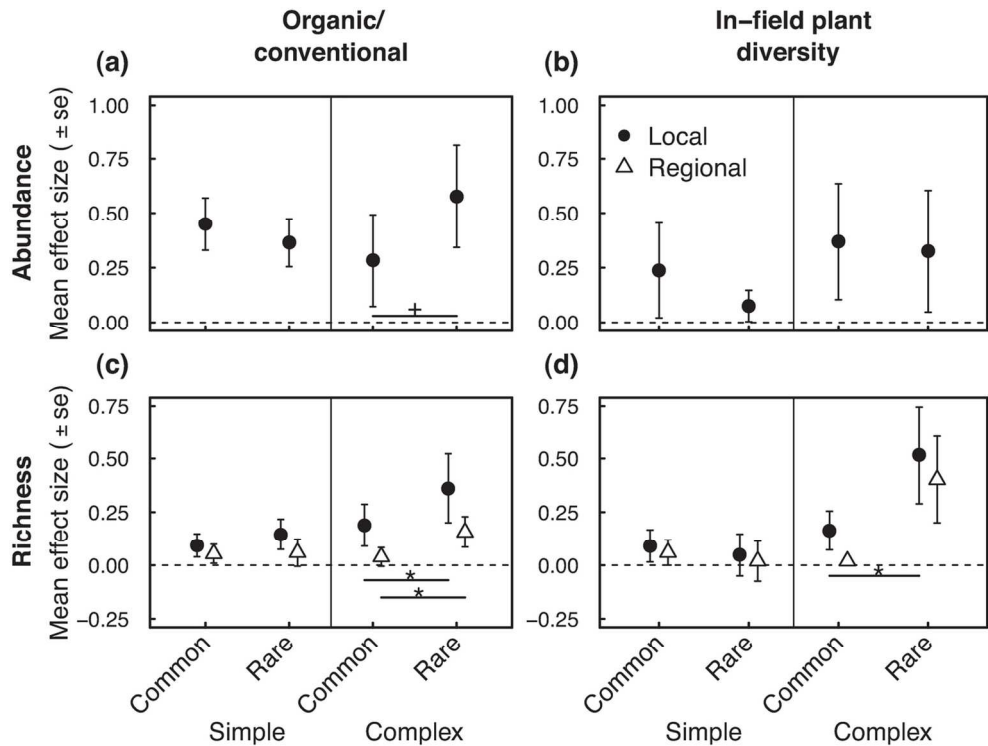


Effects of landscape complexity on the entire arthropod community in organic vs. conventional farms (left column) and fields with high vs. low in-field plant diversity (right column). Each graph shows the mean log-response ratio ( $\pm$  SE) for studies in simple ( $\leq 20\%$  natural habitat) or complex ( $>20\%$  natural habitat) landscapes for (a,b) abundance, (c,d) richness, and (e,f) evenness. A "\*\*\*" ( $p < 0.05$ ) or "+" ( $0.05 \leq p < 0.1$ ) below a set of means indicates a significant difference between means at the habitat complexity levels (Tables S9-S12).

Fig. 3

114x174mm (300 x 300 DPI)





Effects of farm management schemes on abundance (a, b) and richness (c, d) of common vs. rare taxa in simple and complex landscapes. Mean log-response ratios (±SE) of (left column) adopting organic farming and (right column) promoting in-field plant diversity. A "\*" ( $p < 0.05$ ) or "+" ( $0.05 \leq p < 0.1$ ) below a pair of means indicates a significant difference between rare and common taxa within a landscape complexity category (determined via paired t-tests; Table S19).

Fig. 4  
125x93mm (300 x 300 DPI)